

(19) World Intellectual Property
Organization
International Bureau



(43) International Publication Date
29 July 2004 (29.07.2004)

PCT

(10) International Publication Number
WO 2004/062613 A2

(51) International Patent Classification⁷:

A61K

(74) Agents: DUBOFF, Samuel et al.; P.O. Box 4000, Princeton, NJ 08543-4000 (US).

(21) International Application Number:

PCT/US2004/000642

(81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW.

(22) International Filing Date: 12 January 2004 (12.01.2004)

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

60/439,594

13 January 2003 (13.01.2003) US

(71) Applicant (for all designated States except US): BRISTOL-MYERS SQUIBB COMPANY [US/US]; Route 206 And Provinceline Road, Princeton, NJ 08543-4000 (US).

(72) Inventors; and

(75) Inventors/Applicants (for US only): WALKER, Michael, A. [US/US]; 25 Royal Oak Drive, Durham, CT 06422 (US). GULGEZE, Hatice, Belgin [US/US]; 623 Town Colony Drive, Middletown, CT 06457 (US). NAIDU, Narasimhulu, B. [IN/US]; 12 Skeetfield Point Road, Durham, CT 06422 (US). SORENSON, Margaret, E. [US/US]; 27 Harrison Street, Apt. 13, Meriden, CT 06450 (US). UEDA, Yasutsugu [JP/US]; 46 Olde Orchard Road, Clinton, CT 06413 (US). MATISKELLA, John [US/US]; 130 High Hill Road, Wallingford, CT 06492 (US).

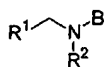
(84) Designated States (unless otherwise indicated, for every kind of regional protection available): ARIPO (BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

Published:

— without international search report and to be republished upon receipt of that report

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: HIV INTEGRASE INHIBITORS



(I)

(57) Abstract: Compounds useful for treating HIV are disclosed having the general formula (I); wherein R¹, R², and B are as defined in the specification. Compositions containing the compounds and methods for using the compounds to inhibit HIV are also disclosed.

WO 2004/062613 A2

BEST AVAILABLE COPY

HIV INTEGRASE INHIBITORS

FIELD OF THE INVENTION

5

The present invention is generally directed to antiviral compounds, and more specifically directed to compounds which can inhibit the functioning of the human immunodeficiency virus-1 (HIV-1), compositions comprising such compounds and methods for inhibiting the functioning of the HIV-1.

10

BACKGROUND OF THE INVENTION

HIV-1 infection remains a major medical problem, with an estimated 42 million people infected worldwide at the end of 2002. The number of cases of HIV and AIDS (acquired immunodeficiency syndrome) has risen rapidly. In 2002, ~5.0 million new infections were reported, and 3.1 million people died from AIDS. Currently available drugs for the treatment of HIV include, for example, the following nucleoside reverse transcriptase (RT) inhibitors: Retrovir™ zidovudine (or AZT), Videx™ didanosine, Zerit™ stavudine, Epivir™ lamivudine (or 3TC), Hivid™ zalcitabine (or DDC), Ziagen™ abacavir succinate, Viread™ Tenofovir disoproxil fumarate salt, Combivir™ (-3TC plus AZT), Trizivir™ (abacavir, lamivudine, and zidovudine); the following non-nucleoside reverse transcriptase inhibitors: Viramune™ nevirapine, Rescriptor™ delavirdine and Sustiva™ efavirenz, and the following peptidomimetic protease inhibitors: saquinavir, indinavir, ritonavir, nelfinavir, amprenavir, lopinavir, Kaletra™ (lopinavir and rtonavir), and Reyataz™ atazanavir sulfate. Often, these drugs can only transiently restrain viral replication if used alone. However, when used in combination, these drugs can have a profound effect on viremia and disease progression. In fact, significant reductions in death rates among AIDS patients have been recently documented as a consequence of the widespread application of combination therapy. However, despite these impressive results, 30 to 50% of patients ultimately fail combination drug therapies. Insufficient drug potency, non-compliance, restricted tissue penetration and drug-specific

limitations within certain cell types (e.g. most nucleoside analogs cannot be phosphorylated in resting cells) may account for the incomplete suppression of sensitive viruses. Furthermore, the high replication rate and rapid turnover of HIV-1 combined with the frequent incorporation of mutations, leads to the appearance of drug-resistant variants and treatment failures when sub-optimal drug concentrations are present. Therefore, novel anti-HIV agents exhibiting distinct resistance patterns, and favorable pharmacokinetic as well as safety profiles are needed to provide more treatment options whether used alone or in combination with other drugs.

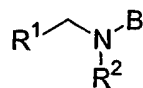
HIV expresses three enzymes, reverse transcriptase, an aspartyl protease and integrase, all of which are potential antiviral targets for the development of drugs for the treatment of AIDS. However, integrase stands out as being the only viral enzyme not targeted by currently approved therapy. The integrase enzyme is responsible for insertion of the viral cDNA into the host cell genome, which is a critical step in the viral life cycle. There are a number of discrete steps involved in this process including processing of the viral cDNA by removal of two bases from each 3'-terminus and joining of the recessed ends to the host DNA. Studies have shown that in the absence of a functional integrase enzyme HIV is not infectious. Therefore, an inhibitor of integrase would be useful as a therapy for AIDS and HIV infection.

A number of HIV integrase inhibitors have been reported. These include nucleotide-based inhibitors, known DNA binders, catechols and hydrazide containing derivatives (Neamati, N.; Sunder, S.; Pommier, Y., *Drug Disc. Today*, 1997, 2, 487).

Certain pyrimidines and pyrimidinones have been disclosed. WO 02/06246 discloses 2-aryl-4,5-dihydroxy-6-carboxypyrimidines as viral polymerase inhibitors which are proposed for use in treating hepatitis C virus infection. Sunderland, C. J.; Botta, M.; Aime, S.; and Raymond, K. N. *Inorg. Chem.* (2001) 40, 6746-6756 discloses the synthesis of 6-carboxamido-5,4-hydroxypyrimidinones as gadolinium chelating agents.

SUMMARY OF INVENTION

The present invention relates to compounds having the formula



(I)

wherein:

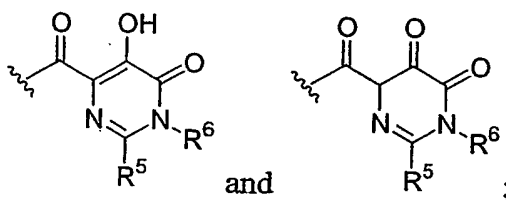
R^1 is $-\text{C}_1\text{--C}_6$ alkyl, $\text{C}_3\text{--C}_6$ cycloalkyl, -aryl, $-\text{C}_1\text{--C}_6$ alkyl-aryl, or $-\text{C}_1\text{--C}_6$ alkyl-diaryl;

R^2 is -H, -OH, $-\text{C}_1\text{--C}_6$ alkyl unsubstituted or substituted with 1-3 R^3 , or $-\text{OR}^4$;

Each R^3 is independently selected from the group consisting of -H, -halo, -CN, $-\text{C}_1\text{--C}_2$ perfluoroalkyl, $-\text{C}_1\text{--C}_6$ alkyl, $-\text{C}_1\text{--C}_2$ perfluoroalkoxy, and $-\text{C}_1\text{--C}_6$ alkoxy;

R^4 is $-\text{C}_1\text{--C}_6$ alkyl or $-\text{C}_3\text{--C}_6$ cycloalkyl;

B is selected from the group consisting of



R^5 is $-\text{NR}^7\text{R}^8$; or Het wherein the bond attaching R^5 is made to a heteroatom in said Het;

R^6 is -H, $-\text{C}_1\text{--C}_6$ alkyl unsubstituted or substituted with 1-3 R^3 ; and

R^7 and R^8 are independently -H or $-\text{C}_1\text{--C}_6$ alkyl;

or pharmaceutically acceptable enantiomer, diastereomer, salt, solvate or prodrug thereof.

The present invention also includes pharmaceutical compositions comprising
5 the compounds or pharmaceutically acceptable enantiomer, diastereomers, salts, solvates or prodrugs thereof and a pharmaceutically acceptable carrier. The compositions can be useful for inhibiting HIV integrase or for treating patients infected with the HIV virus or suffering from AIDS or ARC (AIDS related complex), which comprises a therapeutically effective amount of one or more compounds of
10 Formula I, including pharmaceutical acceptable enantiomer, diastereomers, salts, solvates or prodrugs thereof, and a pharmaceutically acceptable carrier.

The present invention also provides methods of inhibiting HIV integrase by administering to a patient an effective amount of a compound of Formula I, or a
15 pharmaceutically acceptable enantiomer, diastereomers, salt, solvate or prodrug thereof. The present invention further provides methods of treating patients infected by the HIV virus, or of treating AIDS or ARC, by administering to the patient an effective amount of a compound of Formula I, or a pharmaceutically acceptable enantiomer, diastereomers, salt, solvate or prodrug thereof.

20 By virtue of the present invention, it is now possible to provide improved drugs comprising the compounds of the invention which can be effective in the treatment of patients infected with HIV. Specifically, the present invention provides compounds that can inhibit the functioning of HIV integrase. Further, the present
25 invention makes it possible to administer combination therapy to a patient whereby a compound in accordance with the present invention, which is effective to inhibit the HIV integrase, can be administered with another compound having anti-HIV activity, e.g., a compound which is effective to inhibit the function of a target selected from the group consisting of HIV protease, a nucleoside reverse transcription, a non-
30 nucleoside reverse transcriptase or HIV entry.

DETAILED DESCRIPTION OF THE INVENTION

Stereochemical definitions and conventions used herein generally follow McGraw-Hill Dictionary of Chemical Terms, S. P. Parker, Ed., McGraw-Hill Book Company, New York (1984) and Stereochemistry of Organic Compounds, Eliel, E. and Wilen, S., John Wiley & Sons, Inc., New York (1994). Many organic compounds exist in optically active forms, i.e., they have the ability to rotate the plane of plane-polarized light. In describing an optically active compound, the prefixes D and L or R and S are used to denote the absolute configuration of the molecule about its chiral center(s). The prefixes d and l or (+) and (-) are employed to designate the sign of rotation of plane-polarized light by the compound, with (-) or l meaning that the compound is levorotatory and (+) or d, meaning the compound, is dextrorotatory. For a given chemical structure, these compounds, called stereoisomers, are identical except that they are mirror images of one another. A specific stereoisomer of a mirror image pair may also be referred to as an enantiomer, and a mixture of such isomers is often called an enantiomeric mixture. With reference to the instances where (R) or (S) is used, it is to designate the absolute configuration of a substituent in context to the whole compound and not in context to the substituent alone.

20

Unless otherwise specifically noted herein, the terms set forth below will have the following definitions.

The terms "racemic mixture" and "racemate" refer to an equimolar mixture of two enantiomeric species, devoid of optical activity.

25

The term "chiral" refers to molecules which have the property of non-superimposability of the mirror image partner, while the term "achiral" refers to molecules which are superimposable on their mirror image partner.

30

The term "stereoisomers" refers to compounds which have identical chemical composition, but differ with regard to the arrangement of the atoms or groups in space.

5 The term "diastereomer" refers to a stereoisomer which is not an enantiomer, e.g., a stereoisomer with two or more centers of chirality and whose molecules are not mirror images of one another. Diastereomers have different physical properties, e.g. melting points, boiling points, spectral properties, and reactivities. Mixtures of diastereomers may separate under high resolution analytical procedures such as
10 electrophoresis and chromatography.

The term "enantiomers" refers to two stereoisomers of a compound which are non-superimposable mirror images of one another.

15 The term "pharmaceutically acceptable salt" is intended to include nontoxic salts synthesized from a compound which contains a basic or acidic moiety by conventional chemical methods. Generally, such salts can be prepared by reacting the free acid or base forms of these compounds with a stoichiometric amount of the appropriate base or acid in water or in an organic solvent, or in a mixture of the two;
20 generally, nonaqueous media like ether, ethyl acetate, ethanol, isopropanol, or acetonitrile are preferred. Lists of suitable salts are found in *Remington's Pharmaceutical Sciences*, 18th ed., Mack Publishing Company, Easton, PA, 1990, p. 1445. The compounds of the present invention are useful in the form of the free base or acid or in the form of a pharmaceutically acceptable salt thereof. All forms are
25 within the scope of the invention.

The term "therapeutically effective amount" means the total amount of each active component that is sufficient to show a meaningful patient benefit, e.g., a sustained reduction in viral load. When applied to an individual active ingredient,
30 administered alone, the term refers to that ingredient alone. When applied to a combination, the term refers to combined amounts of the active ingredients that result

in the therapeutic effect, whether administered in combination, serially or simultaneously.

The term "compounds of the invention", and equivalent expressions, are meant to embrace compounds of formula I, and pharmaceutically acceptable enantiomers, diastereomers, salts, and solvates, e.g. hydrates, and prodrugs. Similarly, references to intermediates, are meant to embrace their salts, and solvates, where the context so permits. References to the compound of the invention also include the preferred compounds, e.g. formula II and formula III.

10

The term "derivative" means a chemically modified compound wherein the modification is considered routine by the ordinary skilled chemist, such as an ester or an amide of an acid, protecting groups, such as a benzyl group for an alcohol or thiol, and tert-butoxycarbonyl group for an amine.

15

The term "solvate" means a physical association of a compound of this invention with one or more solvent molecules, whether organic or inorganic. This physical association includes hydrogen bonding. In certain instances the solvate will be capable of isolation, for example when one or more solvent molecules are incorporated in the crystal lattice of the crystalline solid. "Solvate" encompasses both solution-phase and isolable solvates. Exemplary solvates include hydrates, ethanولات, methanولات, isopropanولات and the like.

The term "prodrug" as used herein means derivatives of the compounds of the invention which have chemically or metabolically cleavable groups and become, by solvolysis or under physiological conditions, the compounds of the invention which are pharmaceutically active in vivo. A prodrug of a compound may be formed in a conventional manner with a functional group of the compounds such as with an amino, hydroxy or carboxy group when present. The prodrug derivative form often offers advantages of solubility, tissue compatibility, or delayed release in a mammalian organism (see, Bundgard, H., Design of Prodrugs, pp. 7-9, 21-24, Elsevier, Amsterdam 1985). Prodrugs include acid derivatives well known to

30

practitioners of the art, such as, for example, esters prepared by reaction of the parent acidic compound with a suitable alcohol, or amides prepared by reaction of the parent acid compound with a suitable amine.

5 The term "patient" includes both human and other mammals.

 The term "pharmaceutical composition" means a composition comprising a compound of the invention in combination with at least one additional pharmaceutical carrier, i.e., adjuvant, excipient or vehicle, such as diluents,
10 preserving agents, fillers, flow regulating agents, disintegrating agents, wetting agents, emulsifying agents, suspending agents, sweetening agents, flavoring agents, perfuming agents, antibacterial agents, antifungal agents, lubricating agents and dispensing agents, depending on the nature of the mode of administration and dosage forms. Ingredients listed in Remington's Pharmaceutical Sciences, 18th ed., Mack
15 Publishing Company, Easton, PA (1999) for example, may be used.

 The phrase "pharmaceutically acceptable" is employed herein to refer to those compounds, materials, compositions, and/or dosage forms which are, within the scope of sound medical judgment, suitable for use in contact with the tissues of
20 patients without excessive toxicity, irritation, allergic response, or other problem or complication commensurate with a reasonable risk/benefit ratio.

 The term "treating" refers to: (i) preventing a disease, disorder or condition from occurring in a patient which may be predisposed to the disease, disorder and/or
25 condition but has not yet been diagnosed as having it; (ii) inhibiting the disease, disorder or condition, i.e., arresting its development; and (iii) relieving the disease, disorder or condition, i.e., causing regression of the disease, disorder and/or condition.

30 The term "substituted" as used herein includes substitution at from one to the maximum number of possible binding sites on the core, e.p., organic radical, to

which the substituent is bonded, e.g., mono-, di-, tri- or tetra- substituted, unless otherwise specifically stated.

The nomenclature used to describe organic radicals, e.g., hydrocarbons and substituted hydrocarbons, generally follows standard nomenclature known in the art, unless otherwise specifically defined. Combinations of groups, e.g., alkylalkoxyamine or arylalkyl, include all possible stable configurations, unless otherwise specifically stated. Certain radicals and combinations are defined below for purposes of illustration.

10

The term "halo" as used herein means a halogen substituent selected from bromo, chloro, fluoro or iodo. The term "haloalkyl" means an alkyl group that is substituted with one or more halo substituents.

15

The term "alkyl" as used herein means acyclic, straight or branched chain alkyl substituents having the specified number of carbon atoms and includes, for example, methyl, ethyl, propyl, butyl, tert-butyl, hexyl, 1-methylethyl, 1-methylpropyl, 2-methylpropyl, 1,1-dimethylethyl. Thus, C₁₋₆ alkyl refers to an alkyl group having from one to six carbon atoms. The term "lower alkyl" means an alkyl group having from one to six, preferably from one to four carbon atoms. The term "alkylester" means an alkyl group additionally containing an ester group. Generally, a stated carbon number range, e.g., C₂₋₆ alkylester, includes all of the carbon atoms in the radical.

20

The term "alkenyl" as used herein means an alkyl radical containing at least one double bond, e.g., ethenyl (vinyl) and alkyl.

25

The term "alkoxy" as used herein means an alkyl group with the indicated number of carbon atoms attached to an oxygen atom. Alkoxy includes, for example, methoxy, ethoxy, propoxy, 1-methylethoxy, butoxy and 1,1-dimethylethoxy. The latter radical is referred to in the art as tert-butoxy. The term "alkoxycarbonyl" means an alkoxy group additionally containing a carbonyl group.

30

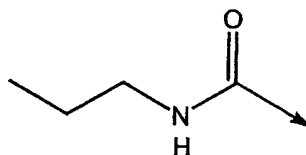
The term "cycloalkyl" as used herein means a cycloalkyl substituent containing the indicated number of carbon atoms and includes, for example, cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl and spiro cyclic groups such as spirocyclopropyl as spirocyclobutyl. The term "cycloalkoxy" as used herein means a cycloalkyl group linked to an oxygen atom, such as, for example, cyclobutyloxy or cyclopropyloxy. The term "alkylcycloalkyl" means a cycloalkyl group linked to an alkyl group. The stated carbon number range includes the total number of carbons in the radical, unless otherwise specifically stated. This a C₄₋₁₀ alkylcycloalkyl may contain from 1-7 carbon atoms in the alkyl group and from 3-9 carbon atoms in the ring, e.g., cyclopropylmethyl or cyclohexylethyl.

The term "aryl" as used herein means an aromatic moiety containing the indicated number of carbon atoms, such as, but not limited to phenyl, indanyl or naphthyl. For example, C₆₋₁₀ aryl refers to an aromatic moiety having from six to ten carbon atoms which may be in the form of a monocyclic or bicyclic structure. The term "haloaryl" as used herein refers to an aryl mono, di or tri substituted with one or more halogen atoms. The terms "alkylaryl", "arylalkyl" and "aralalkyl" mean an aryl group substituted with one or more alkyl groups. Unless the carbon range of each group is specified, the stated range applies to the entire substituent. Thus, a C₇₋₁₄ alkylaryl group may have from 1-8 carbon atoms in the alkyl group for a monocyclic aromatic and from 1-4 carbon atoms in the alkyl group for a fused aromatic. Similarly, a C₁₋₆ alkyl-aryl group would contain a six membered aryl group and a 1-6 carbon alkyl segment. A C₁₋₆ alkyl-diaryl group would contain a ten membered aryl group and a 1-6 carbon alkyl segment. The attachment of the group to bonding site on the molecule can be either at the aryl group or the alkyl group. Unless a specific aryl radical is specified, e.g., fluoro-phenyl, or the radical is stated to be unsubstituted, the aryl radicals include those substituted with typical substituents known to those skilled in the art, e.g., halogen, hydroxy, carboxy, carbonyl, nitro, sulfo, amino, cyano, dialkylamino haloalkyl, CF₃, haloalkoxy, thioalkyl, alkanoyl, SH, alkylamino, alkylamide, dialkylamide, carboxyester,

alkylsulfone, alkylsulfonamide and alkyl(alkoxy)amine. Examples of alkylaryl groups include benzyl, butylphenyl and 1-naphthylmethyl.

The term "alkanoyl" as used herein means straight or branched 1-oxoalkyl radicals containing the indicated number of carbon atoms and includes, for example, 5 formyl, acetyl, 1-oxopropyl (propionyl), 2-methyl-1-oxopropyl, 1-oxohexyl and the like.

The term "alkylamide" as used herein means an amide mono-substituted with 10 an alkyl, such as

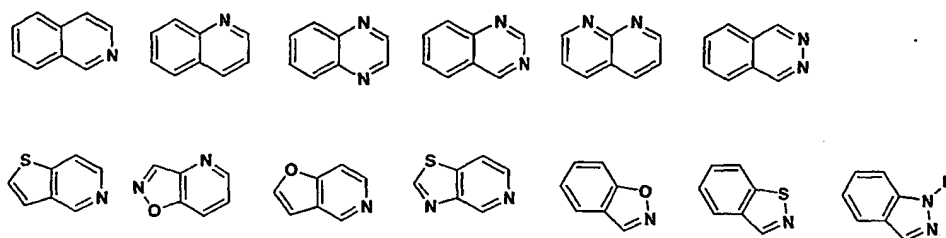


15 The term "heterocycle", also referred to as "Het", as used herein means 7-12 membered bicyclic heterocycles and 5-9 membered monocyclic heterocycles.

Preferred bicyclic heterocycles are 7-12 membered fused bicyclic ring systems (both rings share an adjacent pair of atoms) containing from one to four 20 heteroatoms selected from nitrogen, oxygen and sulfur, wherein one or both rings of the heterocycle can be saturated, partially saturated or fully unsaturated ring system (this latter subset also herein referred to as unsaturated heteroaromatic or heteroaryl). The nitrogen and sulfur heteroatoms atoms may be optionally oxidized. The bicyclic heterocycle may contain the heteroatoms in one or both rings. Unless a specific 25 heterocycle is specified, e.g., a fluorinated 7-12 membered bicyclic heterocycle, or the heterocycle is stated to be unsubstituted, the heterocycles include those substituted with typical substituents known to those skilled in the art. For example, the bicyclic heterocycle may also contain substituents on any of the ring carbon atoms, e.g., one to three substituents. Examples of suitable substituents include C₁₋₆ 30 alkyl, C₃₋₇ cycloalkyl, C₁₋₆ alkoxy, C₃₋₇ cycloalkoxy, halo-C₁₋₆ alkyl, CF₃, mono- or di-

halo- C_{1-6} alkoxy, cyano, halo, thioalkyl, hydroxy, alkanoyl, NO_2 , SH, , amino, C_{1-6} alkylamino, di (C_{1-6}) alkylamino, di (C_{1-6}) alkylamide, carboxyl, (C_{1-6}) carboxyester, C_{1-6} alkylsulfone, C_{1-6} alkylsulfonamide, C_{1-6} alkylsulfoxide, di (C_{1-6}) alkyl(alkoxy)amine, C_{6-10} aryl, C_{7-14} alkylaryl, and a 5-7 membered monocyclic heterocycle. When two substituents are attached to vicinal carbon atoms of the bicyclic heterocycle, they can join to form a ring, e.g., a five, six or seven membered ring system containing up to two heteroatoms selecting from oxygen and nitrogen.

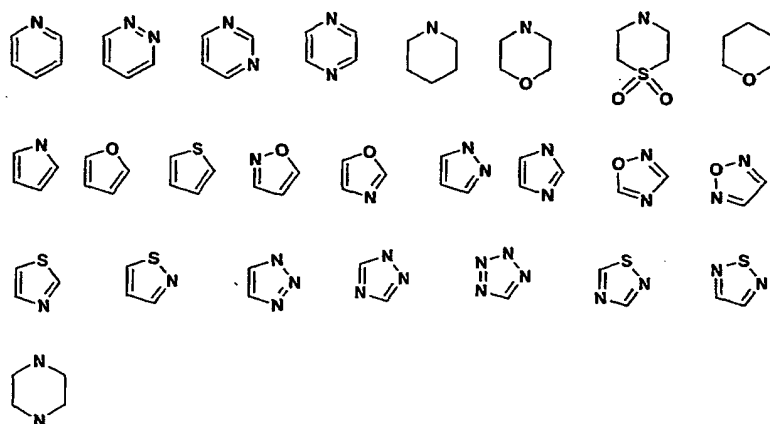
Examples of bicyclic heterocycles include, but are not limited to, the following ring systems:



Preferred monocyclic heterocycles are 5-9 membered saturated, partially saturated or fully unsaturated ring system (this latter subset also herein referred to as unsaturated heteroaromatic) containing in the ring from one to four heteroatoms selected from nitrogen, oxygen and sulfur, wherein the sulfur and nitrogen heteroatoms may be optionally oxidized. Unless a specific heterocycle is specified, e.g., a C_{1-6} alkoxy substituted 5-7 membered monocyclic heterocycle, or the heterocycle is stated to be unsubstituted, the heterocycles include those substituted with typical substituents known to those skilled in the art. For example, the monocyclic heterocycle may also contain substituents on any of the ring atoms, e.g., one to three substituents. Examples of suitable substituents include C_{1-6} alkyl, C_{3-7} cycloalkyl, C_{1-6} alkoxy, C_{3-7} cycloalkoxy, halo- C_{1-6} alkyl, CF_3 , mono-or di- halo- C_{1-6} alkoxy, cyano, halo, thioalkyl, hydroxy, alkanoyl, NO_2 , SH, , amino, C_{1-6} alkylamino, di (C_{1-6}) alkylamino, di (C_{1-6}) alkylamide, carboxyl, (C_{1-6}) carboxyester, C_{1-6} alkylsulfone, C_{1-6} alkylsulfoxide, C_{1-6} alkylsulfonamide, di (C_{1-6}) alkyl(alkoxy)amine, C_{6-10} aryl, C_{7-14} alkylaryl and an additional 5-7 membered monocyclic heterocycle.

Suitable heterocycles include, for example, piperidinyl, piperazinyl, pyrrolidinyl, thiazinyl and morpholinyl. Examples of heteroaryl include 2-furyl, 3-furyl, 2-pyridyl, 3-pyridyl, 4-pyridyl, pyrazinyl, 2-thienyl, 3-thienyl, pyrrolyl, oxazolyl, thiazolyl, imidazolyl, pyrazolyl, isoxazolyl, isothiazolyl, 1,2,3-oxadiazolyl, 1,2,3-triazolyl, 1,3,4-thiadiazolyl, pyridazinyl, pyrimidinyl, 1,3,5-triazinyl and 1,3,5-trithianyl.

Further examples of monocyclic heterocycles include, but are not limited to, the following (and their tautomers):

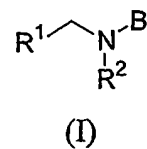


10

Those skilled in the art will recognize that the heterocycles used in the compounds of the present invention should be stable. Generally, stable compounds are those which can be synthesized, isolated and formulated using techniques known to those skilled in the art without degradation of the compound.

15

The present invention provides compounds having the following formula:



20

wherein:

R¹ is -C₁-C₆ alkyl, C₃-C₆ cycloalkyl, -aryl, -C₁-C₆ alkyl-aryl, or -C₁-C₆ alkyl-diaryl;

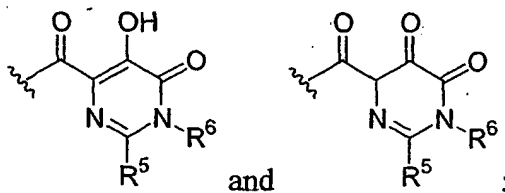
R^2 is -H, -OH, -C₁-C₆ alkyl unsubstituted or substituted with 1-3 R^3 , or -OR⁴;

Each R^3 is independently selected from the group consisting of -H, -halo, -CN, -C₁-
5 C₂ perfluoroalkyl, -C₁-C₆ alkyl, -C₁-C₂ perfluoroalkoxy, and -C₁-C₆ alkoxy;

R^4 is -C₁-C₆ alkyl or -C₃-C₆ cycloalkyl;

B is selected from the group consisting of

10



R^5 is -NR⁷R⁸; or Het wherein the bond attaching R^5 is made to a heteroatom in said
Het;

15

R^6 is -H, -C₁-C₆ alkyl unsubstituted or substituted with 1-3 R^3 ; and

R^7 and R^8 are independently -H or -C₁-C₆ alkyl;

20 or pharmaceutically acceptable enantiomer, diastereomer, salt, solvate or prodrug thereof.

Preferably, the aryl is substituted with 1-3 R^3 . Preferably, R^1 is -aryl, -C₁-C₆ alkyl-aryl, or -C₁-C₆ alkyl diaryl. More preferably, R^1 is -aryl, -C₁-C₆ alkyl-phenyl,
25 or -C₁-C₆ alkyl-diphenyl. Even more preferably, the aryl is selected from the group consisting of phenyl, naphthyl, furanyl, thienyl, indolyl, and benzimidazolyl, and aryl and phenyl are unsubstituted or substituted with 1-3 R^3 . Still more preferably, R^1 is aryl selected from the group consisting of naphthyl, 2-methyl-5-furanyl, 2-thienyl, 3-indolyl, and 2-benzimidazolyl. In a further preferred aspect, R^1 is phenyl

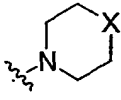
unsubstituted or substituted with 1-3 R³. In a specifically preferred aspect, R¹ is 3-fluorophenyl, 4-fluorophenyl, 2,4-difluorophenyl, 3,4-difluorophenyl, 3-methyl-4-fluorophenyl, 3-chloro-4-fluorophenyl, 3-chlorophenyl, 4-chlorophenyl, 3,4-dichlorophenyl, 3,5-dichlorophenyl, 3-methylphenyl, 4-methylphenyl, or 3,4-dimethylphenyl.

Preferably, R² is -H, methyl, -OH, or methoxy. More preferably, R² is -H.

Preferably, R⁶ is -C₁-C₆ alkyl. More preferably, R⁶ is methyl.

10

Preferably, R⁵ is heteroaryl. More preferably, R⁵ is selected from the group consisting of fluoroindole and methyltriazole. Even more preferably, R⁵ is a 4 to 7 membered monocyclic heterocycle. Still more preferably, R⁵ is selected from the group

consisting of [1,2]thiazinane 1,1-dioxide and , wherein X is a bond, O, S, NH, N-methyl, N-propyl, N-acetyl, N-(2-hydroxyethyl), N-pyridinyl, N-pyrazinyl, N-SO₂Me, CH₂, and CHOH.

Preferred compounds are selected from the group consisting of:

20

N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;

25

N-[(3,4-dichlorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;

30

N-[(3-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;

N-[(4-chlorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;

35

N-[(3-chlorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;

- N-[3-(4-fluorophenyl)propyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 5 N-[3-(3,4-dichlorophenyl)propyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 1,6-dihydro-5-hydroxy-1-methyl-N-[(4-methylphenyl)methyl]-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 10 1,6-dihydro-5-hydroxy-1-methyl-N-[(3-methylphenyl)methyl]-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 1,6-dihydro-5-hydroxy-N-[(4-methoxyphenyl)methyl]-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 15 1,6-dihydro-5-hydroxy-N-[(3-methoxyphenyl)methyl]-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 20 N-[(3,5-dichlorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 25 N-[(3-fluoro-4-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 30 N-[(3,4-difluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- N-[2-(4-fluorophenyl)ethyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 35 N-[(2,4-difluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 40 N-[(3-chloro-4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- N-[(3,4-dimethylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 45 1,6-dihydro-5-hydroxy-1-methyl-6-oxo-N-(1-phenylethyl)-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;

- N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(4-morpholinyl)-6-oxo-4-pyrimidinecarboxamide;
- 5 N-[[4-fluoro-3-(trifluoromethyl)phenyl]methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 10 N-[2-(3-fluorophenyl)ethyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 15 N-[(2,4-dimethoxyphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 1,6-dihydro-5-hydroxy-1-methyl-N-[(5-methyl-2-furanyl)methyl]-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 20 1,6-dihydro-5-hydroxy-N-[(2-methoxyphenyl)methyl]-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- N-(2,2-diphenylethyl)-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 25 N-[2-(4-chlorophenyl)ethyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 1,6-dihydro-5-hydroxy-N-[2-(4-methoxyphenyl)ethyl]-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 30 N-[(2-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-N-[[3-(trifluoromethyl)phenyl]methyl]-4-pyrimidinecarboxamide;
- 35 N-[2-(3-chlorophenyl)ethyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 1,6-dihydro-5-hydroxy-1-methyl-6-oxo-N-(2-phenylethyl)-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 40 1,6-dihydro-5-hydroxy-1-methyl-6-oxo-N-[(1S)-1-phenylethyl]-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 45 1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-N-[[4-(trifluoromethyl)phenyl]methyl]-4-pyrimidinecarboxamide;

- N-[1-(4-fluorophenyl)ethyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 5 N-(2-furanylmethyl)-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-N-(2-thienylmethyl)-4-pyrimidinecarboxamide;
- 10 N-(1,2-diphenylethyl)-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 1,6-dihydro-5-hydroxy-1-methyl-6-oxo-N-(phenylmethyl)-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 15 N-[(2-chlorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 20 1,6-dihydro-5-hydroxy-1-methyl-N-[(2-methylphenyl)methyl]-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 1,6-dihydro-5-hydroxy-N-[2-(2-methoxyphenyl)ethyl]-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 25 1,6-dihydro-5-hydroxy-N-[2-(3-methoxyphenyl)ethyl]-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 1,6-dihydro-5-hydroxy-1-methyl-N-[2-(4-methylphenyl)ethyl]-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 30 1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-N-[(3,4,5-trimethoxyphenyl)methyl]-4-pyrimidinecarboxamide;
- 35 N-[(2,5-dichlorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 1,6-dihydro-5-hydroxy-N-[(1R,2S)-2-hydroxy-1,2-diphenylethyl]-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 40 N-[(2,4-dichlorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 45 N-[[3,5-bis(trifluoromethyl)phenyl]methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;

- N-[(2,5-difluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 5 1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-N-[[2-(trifluoromethyl)phenyl]methyl]-4-pyrimidinecarboxamide;
- 10 1,6-dihydro-5-hydroxy-1-methyl-6-oxo-N-(1-phenylpropyl)-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 15 4-pyrimidinecarboxamide, N-[1-(4-chlorophenyl)ethyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 20 N-[(3,5-difluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 25 1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-N-[[3-(trifluoromethoxy)phenyl]methyl]-4-pyrimidinecarboxamide;
- 30 1,6-dihydro-5-hydroxy-1-methyl-N-[1-(1-naphthalenyl)ethyl]-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 35 N-[1-(4-bromophenyl)ethyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 40 1,6-dihydro-5-hydroxy-1-methyl-N-[1-(2-naphthalenyl)ethyl]-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 45 1,6-dihydro-5-hydroxy-1-methyl-N-[1-(4-methylphenyl)ethyl]-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 1,6-dihydro-5-hydroxy-N-[1-(3-methoxyphenyl)ethyl]-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 1,6-dihydro-5-hydroxy-N-[1-(4-methoxyphenyl)ethyl]-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 1,6-dihydro-5-hydroxy-N-[2-(1H-indol-3-yl)ethyl]-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 1,6-dihydro-5-hydroxy-1-methyl-6-oxo-N-[(1S)-1-phenylpropyl]-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;

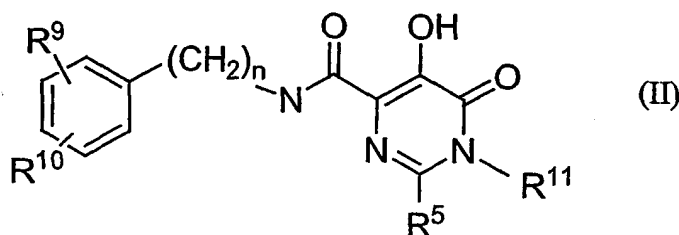
- N-(1H-benzimidazol-2-ylmethyl)-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 5 N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(3-methyl-1H-1,2,4-triazol-1-yl)-6-oxo-4-pyrimidinecarboxamide;
- N-[(3,4-dichlorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(3-methyl-1H-1,2,4-triazol-1-yl)-6-oxo-4-pyrimidinecarboxamide;
- 10 N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(3-methyl-1H-1,2,4-triazol-1-yl)-6-oxo-4-pyrimidinecarboxamide;
- N-[(3,4-dichlorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(4-morpholinyl)-6-oxo-4-pyrimidinecarboxamide;
- 15 N-[(3-fluoro-4-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(4-morpholinyl)-6-oxo-4-pyrimidinecarboxamide;
- N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(4-morpholinyl)-6-oxo-4-pyrimidinecarboxamide;
- 20 N-[(3-chloro-4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(4-morpholinyl)-6-oxo-4-pyrimidinecarboxamide;
- N-[(3,4-dimethylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(4-morpholinyl)-6-oxo-4-pyrimidinecarboxamide;
- 25 N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(5-methyl-1H-1,2,4-triazol-1-yl)-6-oxo-4-pyrimidinecarboxamide;
- 30 N-[(3,4-dichlorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(5-methyl-1H-1,2,4-triazol-1-yl)-6-oxo-4-pyrimidinecarboxamide;
- N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-2-[4-(2-hydroxyethyl)-1-piperazinyl]-1-methyl-6-oxo-4-pyrimidinecarboxamide;
- 35 N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-2-[4-(2-hydroxyethyl)-1-piperazinyl]-1-methyl-6-oxo-4-pyrimidinecarboxamide;
- 40 2-(4-acetyl-1-piperazinyl)-N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-4-pyrimidinecarboxamide;
- 2-(4-acetyl-1-piperazinyl)-N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-4-pyrimidinecarboxamide;
- 45 N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(4-propyl-1-piperazinyl)-4-pyrimidinecarboxamide;

- N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(4-propyl-1-piperazinyl)-4-pyrimidinecarboxamide;
- 5 N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(1-piperazinyl)-4-pyrimidinecarboxamide;
- N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(1-piperazinyl)-4-pyrimidinecarboxamide;
- 10 N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-2-(4-hydroxy-1-piperidinyl)-1-methyl-6-oxo-4-pyrimidinecarboxamide;
- 15 N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-2-(4-hydroxy-1-piperidinyl)-1-methyl-6-oxo-4-pyrimidinecarboxamide;
- 2-(6-fluoro-1,3-dihydro-1-oxo-2H-isoindol-2-yl)-N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-4-pyrimidinecarboxamide;
- 20 N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-[4-(2-pyridinyl)-1-piperazinyl]-4-pyrimidinecarboxamide;
- N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-[4-(2-pyridinyl)-1-piperazinyl]-4-pyrimidinecarboxamide;
- 25 N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-[4-(2-pyrimidinyl)-1-piperazinyl]-4-pyrimidinecarboxamide;
- 30 N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-[4-(2-pyrimidinyl)-1-piperazinyl]-4-pyrimidinecarboxamide;
- N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-[4-(methylsulfonyl)-1-piperazinyl]-6-oxo-4-pyrimidinecarboxamide;
- 35 N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-[4-(methylsulfonyl)-1-piperazinyl]-6-oxo-4-pyrimidinecarboxamide;
- N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(1-pyrrolidinyl)-4-pyrimidinecarboxamide;
- 40 N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-[2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)ethyl]-4-pyrimidinecarboxamide;
- 45 N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-[3-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)propyl]-4-pyrimidinecarboxamide;

N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-[2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)ethyl]-4-pyrimidinecarboxamide;
and

N-[(3,4-dichlorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-[2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)ethyl]-4-pyrimidinecarboxamide.

In another aspect of the invention, there is provided a compound having the formula:



wherein,

n is 1 to 4;

R⁵ is Het wherein the bond attaching R⁵ is made to a heteroatom in said Het;

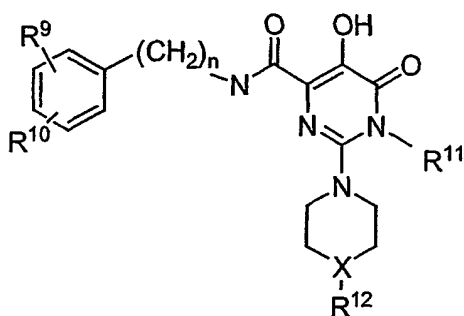
R⁹ and R¹⁰ are each independently H, halo, C₁₋₆ alkyl, C₁₋₆ alkoxy or CF₃; and

R¹¹ is C₁₋₆ alkyl;

or a pharmaceutically acceptable enantiomer, diastereomer, salt, solvate or prodrug thereof.

Preferably, R⁵ is a C₄₋₇ monocyclic heterocycle. More preferably, the heterocycle is substituted with at least one of C₁₋₆ alkyl, C₁₋₆ alkoxy, C₁₋₆ hydroxyalkyl, C₁₋₆ alkyl carbonyl, C₆₋₁₀ aryl, C₇₋₁₄ alkyl, aryl, C₁₋₆ alkylsulfonyl, C₆₋₁₀ arylsulfonyl, C₁₋₆ alkylurea, C₃₋₇ cycloalkylurea, carbonyl, carboxamide or hydroxyl.

In another aspect of the invention, there is provided a compound having the formula:



(III)

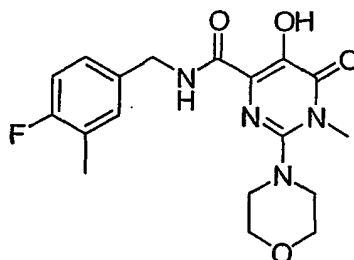
- 5 wherein n is 1 to 4;
 R^9 and R^{10} are each independently H, halo, C_{1-6} alkyl, C_{1-6} alkoxy or CF_3 ;
 X is C, O or N;
 R^{11} is C_{1-6} alkyl; and
 R^{12} is H, C_{1-6} alkyl, C_{1-6} alkoxy, C_{1-6} hydroxyalkyl, C_{1-6} alkyl carbonyl, C_{6-10} aryl, C_{7-14} alkyl; aryl, C_{1-6} alkylsulfonyl, C_{6-10} arylsulfonyl, C_{1-6} alkylurea, C_{3-7} cycloalkylurea,
 10 carbonyl, carboxamide or hydroxyl;
 said R^{12} being present only when X is C or N;
 or a pharmaceutically acceptable enantiomer, diastereomer, salt, solvate or prodrug thereof.

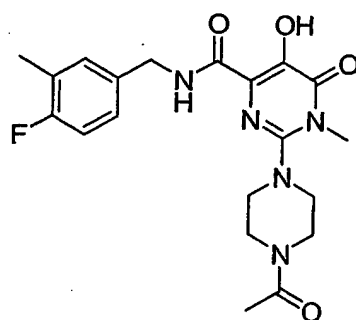
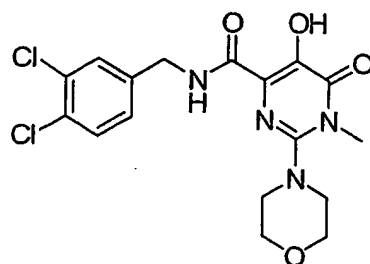
15

Preferably, n is 1. Preferably, R_9 is halo or methyl. More preferably, R_9 is fluoro or chloro. Preferably, R_{10} is methyl. More preferably R_{10} is chloro. Preferably, R_9 and R_{10} are not both H. Preferably, X is O or N.

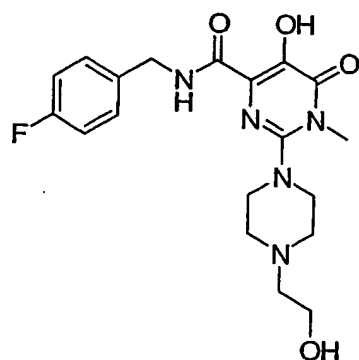
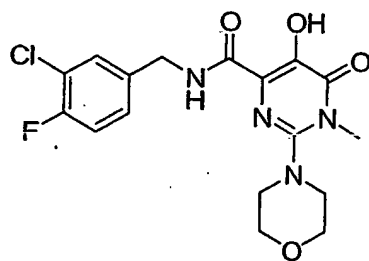
20

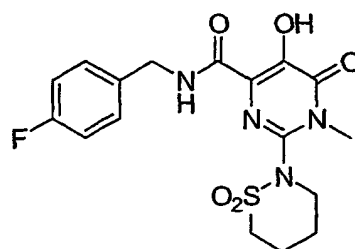
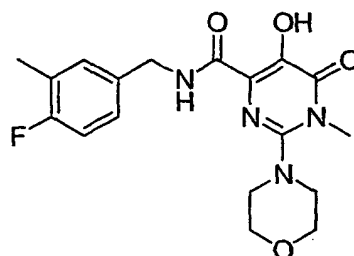
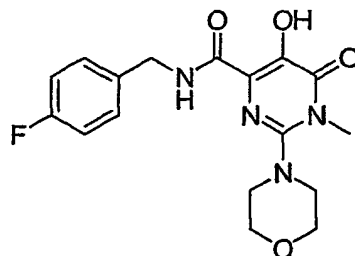
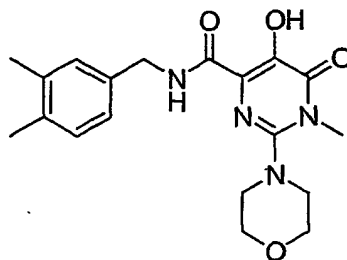
Preferred compounds of the invention have the following structures:





5





5

10 The compounds of the present invention, which are substituted with a basic group, can form salts by the addition of a pharmaceutically acceptable acid. The acid addition salts are formed from a compound of Formula I and a pharmaceutically acceptable inorganic acid, including but not limited to hydrochloric, hydrobromic, hydroiodic, sulfuric, phosphoric, or organic acid such as *p*-toluenesulfonic, methanesulfonic, acetic, benzoic, citric, malonic, fumaric, maleic, oxalic, succinic,

15

sulfamic, or tartaric. Thus, examples of such pharmaceutically acceptable salts include chloride, bromide, iodide, sulfate, phosphate, methanesulfonate, citrate, acetate, malonate, fumarate, sulfamate, and tartrate.

5 Salts of an amine group may also comprise quaternary ammonium salts in which the amino nitrogen carries a suitable organic group such as an alkyl, alkenyl, alkynyl or aralkyl moiety.

 Compounds of the present invention, which are substituted with an acidic
10 group, may exist as salts formed through base addition. Such base addition salts include those derived from inorganic bases which include, for example, alkali metal salts (e.g. sodium and potassium), alkaline earth metal salts (e.g. calcium and magnesium), aluminum salts and ammonium salts. In addition, suitable base addition salts include salts of physiologically acceptable organic bases such as
15 trimethylamine, triethylamine, morpholine, pyridine, piperidine, picoline, dicyclohexylamine, N,N'-dibenzylethylenediamine, 2-hydroxyethylamine, bis-(2-hydroxyethyl)amine, tri-(2-hydroxyethyl)amine, procaine, dibenzylpiperidine, N-benzyl- β -phenethylamine, dehydroabietylamine, N,N'-bishydroabietylamine, glucamine, N-methylglucamine, collidine, quinine, quinoline, ethylenediamine,
20 ornithine, choline, N,N'-benzylphenethylamine, chlorprocaine, diethanolamine, diethylamine, piperazine, tris(hydroxymethyl)aminomethane and tetramethylammonium hydroxide and basic amino acids such as lysine, arginine and N-methylglutamine. These salts may be prepared by methods known to those skilled in the art.

25

 Certain compounds of the present invention, and their salts, may also exist in the form of solvates with water, for example hydrates, or with organic solvents such as methanol, ethanol or acetonitrile to form, respectively, a methanolate, ethanolate or acetonitrilate. The present invention includes each solvate and mixtures thereof.

30

In addition, compounds of the present invention, or a salt or solvate thereof, may exhibit polymorphism. The present invention also encompasses any such polymorphic form.

5 The present invention includes both enantiomers and mixtures of enantiomers such as racemic mixtures.

 The enantiomers may be resolved by methods known to those skilled in the art, for example, by formation of diastereoisomeric salts which may be separated by
10 crystallization, gas-liquid or liquid chromatography, selective reaction of one enantiomer with an enantiomer-specific reagent. It will be appreciated that where the desired enantiomer is converted into another chemical entity by a separation technique, then an additional step is required to form the desired enantiomeric form. Alternatively, specific enantiomers may be synthesized by asymmetric synthesis
15 using optically active reagents, substrates, catalysts or solvents, or by converting one enantiomer into the other by asymmetric transformation.

 The compounds of the present invention may be in the form of a prodrug. Simple aliphatic or aromatic esters derived from, when present, acidic groups
20 pendent on the compounds of this invention are preferred prodrugs. In some cases it is desirable to prepare double ester type prodrugs such as (acyloxy) alkyl esters or (alkoxycarbonyl)oxy)alkyl esters.

 Certain compounds of the present invention may also exist in different stable
25 conformational forms which may be separable. Torsional asymmetry due to restricted rotation about an asymmetric single bond, for example because of steric hindrance or ring strain, may permit separation of different conformers. The present invention includes each conformational isomer of these compounds and mixtures thereof.

Certain compounds of the present invention may exist in zwitterionic form and the present invention includes each zwitterionic form of these compounds and mixtures thereof.

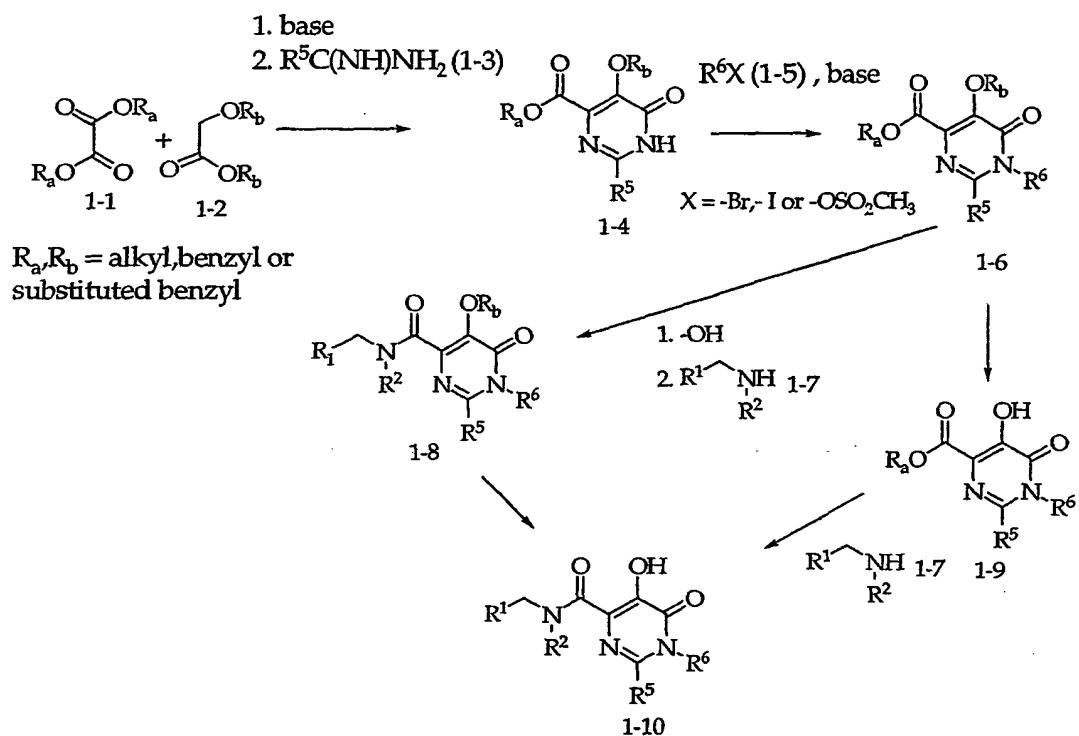
5 The starting materials useful to synthesize the compounds of the present invention are known to those skilled in the art and can be readily manufactured or are commercially available.

10 The compounds of the present invention can be manufactured by methods known to those skilled in the art, see e.g., (Sunderland, J. S.; Botta, M.; Aime, S.; Raymond, K. N. *Inorg. Chem.* (2001), 40, 6756-6756). The following methods set forth below are provided for illustrative purposes and are not intended to limit the scope of the claimed invention. It will be recognized that it may be preferred or necessary to prepare such a compound in which a functional group is protected using
15 a conventional protecting group then to remove the protecting group to provide a compound of the present invention. The details concerning the use of protecting groups in accordance with the present invention are known to those skilled in the art.

20 The compounds of the present invention can be synthesized according to Scheme I. In Scheme I, an oxalic acid diester 1-1 is condensed with glycolate 1-2 using sodium hydride or a similar base. The intermediate generated from this reaction can be isolated but more often is reacted in one pot with an appropriately substituted amidine (1-3) to yield the pyrimidinone heterocycle 1-4. This is alkylated with a suitable electrophile, 1-5, under basic conditions. From here there are two
25 alternative pathways to the final product. In one path intermediate 1-6 is treated with base to saponify the ester and the resulting acid coupled with amine 1-7. It will be appreciated by those skilled in the art that the amide coupling reaction can be carried out under a variety of conditions such as those disclosed in Jerry March, *Advanced Organic Chemistry*, 3rd edition, John Wiley & Sons, 1985. The resulting amide, 1-8
30 is then treated under conditions appropriate for cleaving the protecting group R_b to yield 1-10. Where R_b is alkyl, this can be accomplished by BBr₃ or other conditions familiar to those skilled in the art, such as treatment with LiI. Alternatively when R_b

is a benzylic or substituted benzylic group the ether can be cleaved under reductive conditions, oxidative conditions or acidic conditions. Protecting groups, R, useful for the synthesis of compounds such as 1-9 can be found in Greene, T. W. and Wutz, P. G. M. Protective Groups in Organic Synthesis, Second Edition, 1991, John Wiley and Sons, New York. In the alternative pathway, the protecting group is removed from 1-6 to yield 1-9. At elevated temperatures 1-9 reacts with amines to afford 1-10.

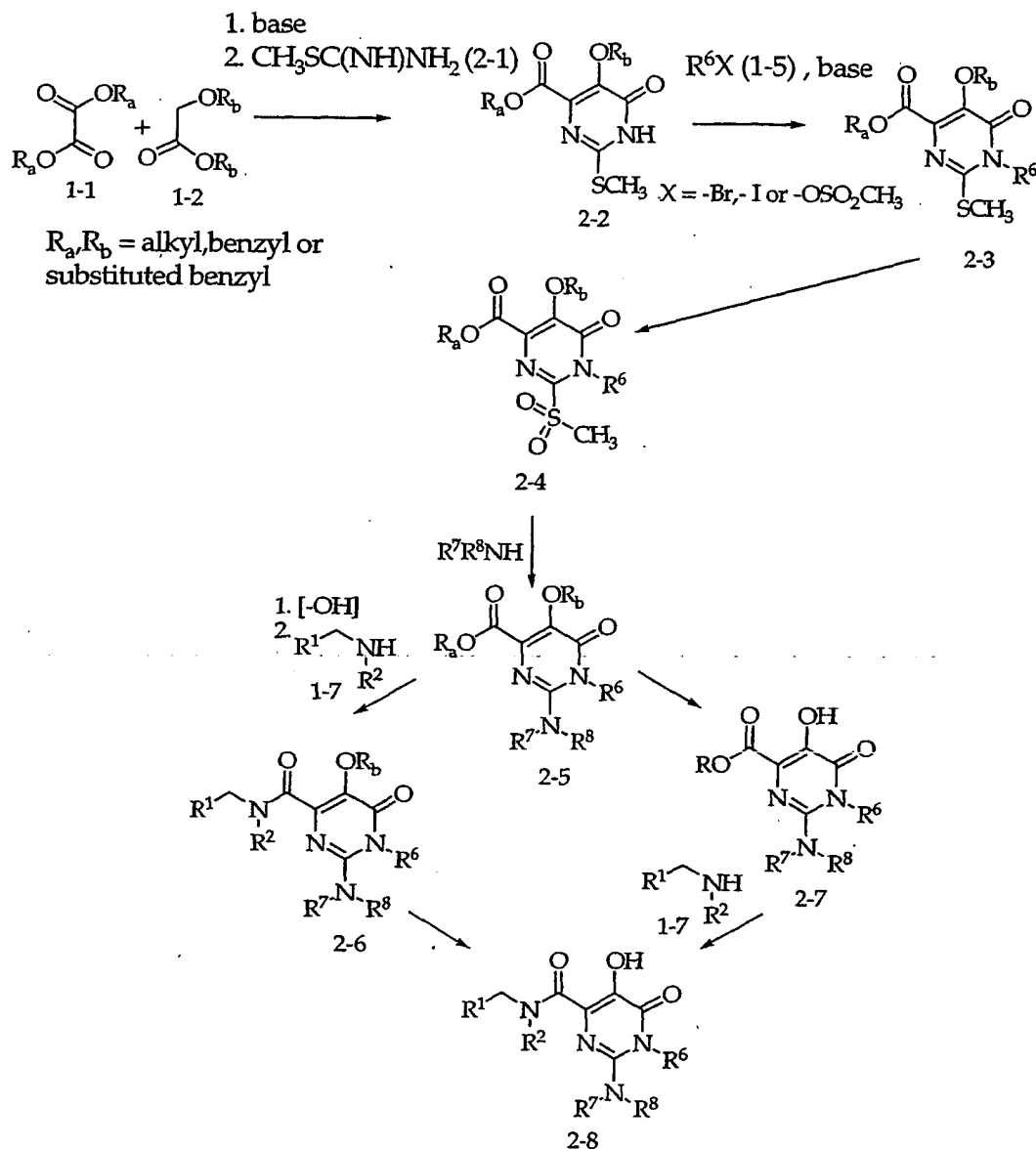
Scheme I



In Scheme II, an alternative pathway is shown in which the R⁵ group is introduced at a later stage of the synthesis. Thus, 2-2 is formed from an oxalate ester, protected glycolic acid and S-methyl-thiourea using the same conditions as described in Scheme I. Likewise, this intermediate is carried through the same sequence of reactions described in Scheme I to yield 2-4. The thiomethyl group is then displaced with a nitrogen containing heterocycle, such as those shown in the scheme, under acidic conditions (HCl) similar to those described in Ple, N.; Turck, Heynderick, A.; Queguiner, G. J. Heterocycl. Chem. (1997) 34, 551-556, for example, or after oxidation of the sulphur atom as described in Jacobs, R. T.; Mauger, R. C.; Ulatowski, T. G.; Aharony, D.; Buckner, C. K. Bioorg. Med. Chem. Lett. (1995), 23, 2879-2884. Following the installation of the amine at C2 of the pyrimidine template

there are two alternative pathways for introducing the amide group. These are essentially the same as those described in Scheme I

Scheme II



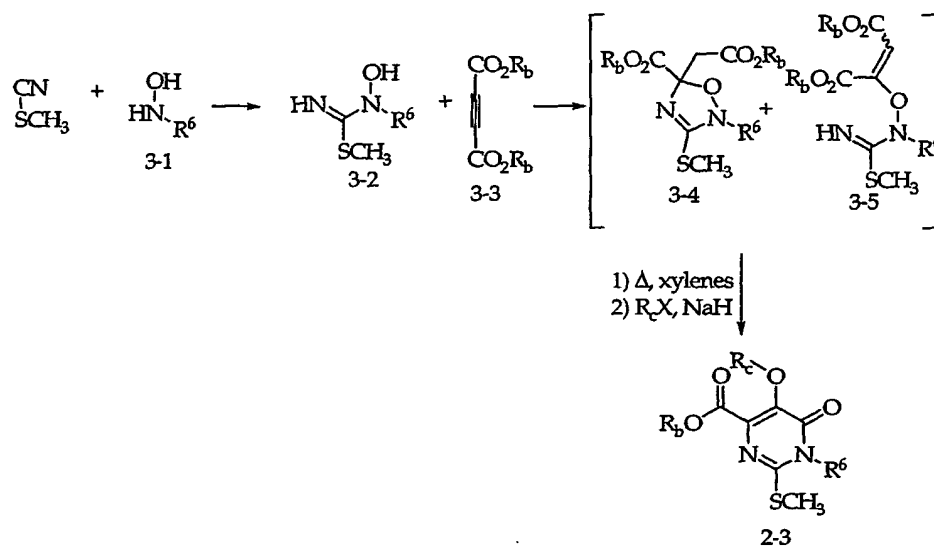
5

A different route to intermediate 2-3 is shown in scheme III.

Methylthiocyanate reacts with appropriately substituted amines to form N-hydroxyl amidines such as 3-2. This intermediate will react with an alkyne-diester to afford a mixture of 3-4 and 3-5. Both of these compounds will rearrange to form the hydroxypyrimidinone template (Culbertson, T. P.; J. Heterocyclic Chem. (1979) 16, 1423-1424.). The in-situ reaction with an appropriate electrophile results in 2-3.

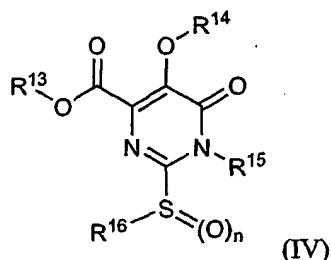
10

Scheme III



- 5 The compounds of the present invention can, for example, be preferably synthesized according to Scheme I. In Scheme I, an oxalic acid diester 1-1 is condensed with glycolate 1-2 using sodium hydride or a similar base. The intermediate generated from this reaction can be isolated but more often is reacted in one pot with an appropriately substituted amidine (1-3) to yield the pyrimidinone
- 10 heterocycle 1-4. This is alkylated with a suitable electrophile, 1-5, under basic conditions. Intermediate 1-6 is treated with base to saponify the ester and the resulting acid coupled with amine 1-7. It will be appreciated by those skilled in the art that the amide coupling reaction can be carried out under a variety of conditions such as those disclosed in Jerry March, *Advanced Organic Chemistry*, 3rd edition,
- 15 John Wiley & Sons, 1985. The resulting amide, 1-8 is then treated under conditions appropriate for cleaving the protecting group R. For alkyl groups, where R is alkyl, this can be accomplished by BBr_3 or other conditions familiar to those skilled in the art, such as treatment with LiI. Alternatively when R is a benzylic or substituted benzylic group the ether can be cleaved under reductive conditions, oxidative
- 20 conditions or acidic conditions. Protecting groups, R, useful for the synthesis of compounds such as 1-9 can be found in Greene, T. W. and Wutz, P. G. M. *Protective Groups in Organic Synthesis*, Second Edition, 1991, John Wiley and Sons, New York.

A preferred intermediate for use in accordance with the present invention is a compound of the formula:



5

wherein;

10- n is 0, 1 or 2;

R^{13} is -H, -C₁₋₆ alkyl, C₁₋₆ cycloalkyl, -aryl, C₁₋₆ alkyl-cycloalkyl, -C₁₋₆ alkyl-aryl, or -C₁₋₆ alkyl-diaryl,

15 R^{14} is -H, -C₁₋₆ alkyl, C₁₋₆ cycloalkyl, -aryl, C₁₋₆ alkyl-aryl, or -C₁₋₆ alkyl-diaryl,

R^{15} is -H, -C₁₋₆ alkyl, C₁₋₆ alkyl-cycloalkyl, -aryl, or -C₁₋₆ alkyl-aryl, and

R^{16} is -H, -C₁₋₆ alkyl, C₁₋₆ cycloalkyl, -aryl, C₁₋₆ alkyl-aryl, or -C₁₋₆ alkyl-diaryl.

20

The present invention also provides compositions comprising a compound of the present invention, or a pharmaceutically acceptable enantiomer, diastereomer, salt, solvate or prodrug thereof, and a pharmaceutically acceptable carrier.

25 Pharmaceutical compositions of the present invention comprise a therapeutically effective amount of a compound of the invention, or a pharmaceutically acceptable enantiomer, diastereomer, salt, solvate or prodrug thereof, and a pharmaceutically acceptable carrier, with a pharmaceutically acceptable carrier, e.g., excipient, or vehicle diluent.

30

The active ingredient, i.e., compound, in such compositions typically comprises from 0.1 weight percent to 99.9 percent by weight of the composition, and often comprises from about 5 to 95 weight percent.

5 Thus, in one aspect of the invention, there is provided a composition comprising the compound of formula 1 and a pharmaceutically acceptable carrier. Preferably, the composition further comprises a compound having anti-HIV activity. As used herein, the term "anti-HIV activity" means the compound is effective to inhibit the function of a target selected from the group consisting of HIV protease, a
10 nucleoside reverse transcriptase, a non-nucleoside reverse transcriptase or HIV entry. Often, the other compound having anti-HIV activity is effective to inhibit the function of target in the HIV life cycle other than HIV integrase.

In a preferred aspect of the invention, the compound having anti-HIV activity
15 is a small molecule compound. As used herein, the term "small molecule compound" means a compound having a molecular weight of less than 1,500 daltons, preferably less than 1000 daltons. Preferably, the small molecule compound is effective to inhibit the function of a target selected from the group consisting of HIV protease, a nucleoside reverse transcriptase, a non-nucleoside reverse transcriptase or HIV entry.

20

Preferred compounds having anti-HIV activity include those selected from the group consisting of HIV protease inhibitors, nucleoside reverse transcriptase inhibitors, non-nucleoside reverse transcriptase inhibitors, HIV-entry inhibitors, immunomodulators, antiinfectives, and vaccines.

25

Thus the present invention is also directed to combinations of the compounds with one or more agents useful in the treatment of AIDS. For example, the compounds of this invention may be effectively administered, whether at periods of pre-exposure and/or post-exposure, in combination with effective amounts of the
30 AIDS antivirals, immunomodulators, antiinfectives, or vaccines, such as those in the following table.

TABLE 1
ANTIVIRALS

	<u>Drug Name</u>	<u>Manufacturer</u>	<u>Indication</u>
5	097	Hoechst/Bayer	HIV infection, AIDS, ARC (non-nucleoside reverse trans- criptase (RT) inhibitor)
10			
15	Amprenivir 141 W94 GW 141	Glaxo Wellcome	HIV infection, AIDS, ARC (protease inhibitor)
20	Abacavir (1592U89) GW 1592	Glaxo Wellcome	HIV infection, AIDS, ARC (RT inhibitor)
	Acemannan	Carrington Labs (Irving, TX)	ARC
25	Acyclovir	Burroughs Wellcome	HIV infection, AIDS, ARC, in combination with AZT
30	AD-439	Tanox Biosystems	HIV infection, AIDS, ARC
	AD-519	Tanox Biosystems	HIV infection, AIDS, ARC
35	Adefovir dipivoxil AL-721	Gilead Sciences Ethigen (Los Angeles, CA)	HIV infection ARC, PGL HIV positive, AIDS
	Alpha Interferon	Glaxo Wellcome	Kaposi's sarcoma,

			HIV in combination w/Retrovir
5	Ansamycin LM 427	Adria Laboratories (Dublin, OH) Erbamont (Stamford, CT)	ARC
10	Antibody which Neutralizes pH Labile alpha aberrant Interferon	Advanced Biotherapy Concepts (Rockville, MD)	AIDS, ARC
15	AR177	Aronex Pharm	HIV infection, AIDS, ARC
	Beta-fluoro-ddA	Nat'l Cancer Institute	AIDS-associated diseases
20	Reyataz™ atazanavir sulfate	Bristol-Myers Squibb/ Novartis	HIV infection, AIDS, ARC (protease inhibitor)
25	CI-1012	Warner-Lambert	HIV-1 infection
	Cidofovir	Gilead Science	CMV retinitis, herpes, papillomavirus
30	Curdlan sulfate	AJI Pharma USA	HIV infection
	Cytomegalovirus Immune globin	MedImmune	CMV retinitis
35	Cytovene	Syntex	Sight threatening
	Ganciclovir		CMV peripheral CMV retinitis
40	Delaviridine	Pharmacia-Upjohn	HIV infection,

			AIDS, ARC (RT inhibitor)
5	Dextran Sulfate	Ueno Fine Chem. Ind. Ltd. (Osaka, Japan)	AIDS, ARC, HIV positive asymptomatic
10	ddC Dideoxycytidine	Hoffman-La Roche	HIV infection, AIDS, ARC
	ddI Dideoxyinosine	Bristol-Myers Squibb	HIV infection, AIDS, ARC; combination with AZT/d4T
15	DMP-450	AVID (Camden, NJ)	HIV infection, AIDS, ARC (protease inhibitor)
20	Efavirenz (-)-6-Chloro-4-(S)- cyclopropylethynyl- 4(S)-trifluoro- methyl-1,4-dihydro- 2H-3,1-benzoxazin- 25 2-one, STOCRINE	DuPont Merck	HIV infection, AIDS, ARC (non-nucleoside RT inhibitor)
	EL10	Elan Corp, PLC (Gainesville, GA)	HIV infection
30	Famciclovir	Smith Kline	herpes zoster, herpes simplex
35	FTC	Emory University	HIV infection, AIDS, ARC (reverse transcriptase inhibitor)
40	GS 840	Gilead	HIV infection, AIDS, ARC (reverse transcriptase inhibitor)

5	HBV097	Hoechst Marion Roussel	HIV infection, AIDS, ARC (non-nucleoside reverse transcriptase inhibitor)
	Hypericin	VIMRx Pharm.	HIV infection, AIDS, ARC
10	Recombinant Human Interferon Beta	Triton Biosciences (Alameda, CA)	AIDS, Kaposi's sarcoma, ARC
	Interferon alfa-n3	Interferon Sciences	ARC, AIDS
15	Indinavir	Merck	HIV infection, AIDS, ARC, asymptomatic HIV positive, also in combination with AZT/ddI/ddC
20	ISIS 2922	ISIS Pharmaceuticals	CMV retinitis
	KNI-272	Nat'l Cancer Institute	HIV-assoc. diseases
25	Lamivudine, 3TC	Glaxo Wellcome	HIV infection, AIDS, ARC (reverse transcriptase inhibitor); also with AZT
30	Lobucavir	Bristol-Myers Squibb	CMV infection
35	Nelfinavir	Agouron Pharmaceuticals	HIV infection, AIDS, ARC (protease inhibitor)
	Nevirapine	Boehringer Ingelheim	HIV infection, AIDS, ARC (RT inhibitor)
40			

	Novapren	Novaferon Labs, Inc. (Akron, OH)	HIV inhibitor
5	Peptide T Octapeptide Sequence	Peninsula Labs (Belmont, CA)	AIDS
10	Trisodium Phosphonoformate	Astra Pharm. Products, Inc.	CMV retinitis, HIV infection, other CMV infections
15	PNU-140690	Pharmacia Upjohn	HIV infection, AIDS, ARC (protease inhibitor)
	Probucol	Vyrex	HIV infection, AIDS
20	RBC-CD4	Sheffield Med. Tech (Houston, TX)	HIV infection, AIDS, ARC
	Ritonavir	Abbott	HIV infection, AIDS, ARC (protease inhibitor)
25	Saquinavir	Hoffmann- LaRoche	HIV infection, AIDS, ARC (protease inhibitor)
30	Stavudine; d4T Didehydrodeoxy- thymidine	Bristol-Myers Squibb	HIV infection, AIDS, ARC
35	Valaciclovir	Glaxo Wellcome	Genital HSV & CMV infections
	Virazole Ribavirin	Viratek/ICN (Costa Mesa, CA)	asymptomatic HIV positive, LAS, ARC
40	VX-478	Vertex	HIV infection, AIDS, ARC

	Zalcitabine	Hoffmann-LaRoche	HIV infection, AIDS, ARC, with AZT
5	Zidovudine; AZT	Glaxo Wellcome	HIV infection, AIDS, ARC, Kaposi's sarcoma, in combination with other therapies
10	Tenofovir disoproxil, fumarate salt (Viread [®])	Gilead	HIV infection, AIDS, (reverse transcriptase inhibitor)
15	Combivir [®]	GSK	HIV infection, AIDS, (reverse transcriptase inhibitor)
20	abacavir succinate (or Ziagen [®])	GSK	HIV infection, AIDS, (reverse transcriptase inhibitor)
25	Fuzeon [®] (or T-20)	Roche / Trimeris AIDs, viral Fusion	HIV infection inhibitor

IMMUNOMODULATORS

30	<u>Drug Name</u>	<u>Manufacturer</u>	<u>Indication</u>
	AS-101	Wyeth-Ayerst	AIDS
	Bropirimine	Pharmacia Upjohn	Advanced AIDS
35	Acemannan	Carrington Labs, Inc. (Irving, TX)	AIDS, ARC
	CL246,738	American Cyanamid	AIDS, Kaposi's

		Lederle Labs	sarcoma
	EL10	Elan Corp, PLC (Gainesville, GA)	HIV infection
5	FP-21399	Fuki ImmunoPharm	Blocks HIV fusion with CD4+ cells
10	Gamma Interferon	Genentech	ARC, in combination w/TNF (tumor necrosis factor)
15	Granulocyte Macrophage Colony Stimulating Factor	Genetics Institute Sandoz	AIDS
20	Granulocyte Macrophage Colony Stimulating Factor	Hoechst-Roussel Immunex	AIDS
	Granulocyte Macrophage Colony Stimulating Factor	Schering-Plough	AIDS, combination w/AZT
25	HIV Core Particle Immunostimulant	Rorer	Seropositive HIV
30	IL-2 Interleukin-2	Cetus	AIDS, in combination w/AZT
	IL-2 Interleukin-2	Hoffman-LaRoche Immunex	AIDS, ARC, HIV, in combination w/AZT
35	IL-2 Interleukin-2 (aldeshukin)	Chiron	AIDS, increase in CD4 cell counts

	Immune Globulin Intravenous (human)	Cutter Biological (Berkeley, CA)	Pediatric AIDS, in combination w/AZT
5	IMREG-1	Imreg (New Orleans, LA)	AIDS, Kaposi's sarcoma, ARC, PGL
	IMREG-2	Imreg (New Orleans, LA)	AIDS, Kaposi's sarcoma, ARC, PGL
10	Imuthiol Diethyl Dithio Carbamate	Merieux Institute	AIDS, ARC
	Alpha-2	Schering Plough	Kaposi's sarcoma
15	Interferon		w/AZT, AIDS
	Methionine- Enkephalin	TNI Pharmaceutical (Chicago, IL)	AIDS, ARC
20	MTP-PE Muramyl-Tripeptide	Ciba-Geigy Corp.	Kaposi's sarcoma
	Granulocyte Colony Stimulating Factor	Amgen	AIDS, in combination w/AZT
25	Remune	Immune Response Corp.	Immunotherapeutic
30	rCD4 Recombinant Soluble Human CD4	Genentech	AIDS, ARC
35	rCD4-IgG hybrids		AIDS, ARC
40	Recombinant Soluble Human CD4	Biogen	AIDS, ARC

	Interferon Alfa 2a	Hoffman-La Roche	Kaposi's sarcoma AIDS, ARC, in combination w/AZT
5	SK&F106528 Soluble T4	Smith Kline	HIV infection
10	Thymopentin	Immunobiology Research Institute (Annandale, NJ)	HIV infection
	Tumor Necrosis Factor; TNF	Genentech	ARC, in combination w/gamma Interferon

15

ANTI-INFECTIVES

	<u>Drug Name</u>	<u>Manufacturer</u>	<u>Indication</u>
20	Clindamycin-with Primaquine	Pharmacia Upjohn	PCP
25	Fluconazole	Pfizer	Cryptococcal meningitis, candidiasis
	Pastille Nystatin Pastille	Squibb Corp.	Prevention of oral candidiasis
30	Ornidyl Eflornithine	Merrell Dow	PCP
	Pentamidine Isethionate (IM & IV)	LyphoMed (Rosemont, IL)	PCP treatment
35	Trimethoprim		Antibacterial
	Trimethoprim/sulfa		Antibacterial
	Piritrexim	Burroughs Wellcome	PCP treatment

5	Pentamidine Isethionate for Inhalation	Fisons Corporation	PCP prophylaxis
	Spiramycin	Rhone-Poulenc diarrhea	Cryptosporidial
10	Intraconazole- R51211	Janssen-Pharm.	Histoplasmosis; cryptococcal meningitis
	Trimetrexate	Warner-Lambert	PCP
15	Daunorubicin	NeXstar, Sequus	Kaposi's sarcoma
	Recombinant Human Erythropoietin	Ortho Pharm. Corp.	Severe anemia assoc. with AZT therapy
20	Recombinant Human Growth Hormone	Serono	AIDS-related wasting, cachexia
25	Megestrol Acetate	Bristol-Myers Squibb	Treatment of anorexia assoc. W/AIDS
	Testosterone	Alza, Smith Kline	AIDS-related wasting
30	Total Enteral Nutrition	Norwich Eaton Pharmaceuticals	Diarrhea and malabsorption related to AIDS

Additionally, the compounds of the invention herein may be used in
 35 combination with another class of agents for treating AIDS which are called HIV
 entry inhibitors. Examples of such HIV entry inhibitors are discussed in DRUGS OF
 THE FUTURE 1999, 24(12), pp. 1355-1362; CELL, Vol. 9, pp. 243-246, Oct. 29,
 1999; and DRUG DISCOVERY TODAY, Vol. 5, No. 5, May 2000, pp. 183-194.

It will be understood that the scope of combinations of the compounds of this invention with AIDS antivirals, immunomodulators, anti-infectives, HIV entry inhibitors or vaccines is not limited to the list in the above Table 1, but includes in principle any combination with any pharmaceutical composition useful for the treatment of AIDS.

Preferred combinations are simultaneous or alternating treatments of with a compound of the present invention and an inhibitor of HIV protease and/or a non-nucleoside inhibitor of HIV reverse transcriptase. An optional fourth component in the combination is a nucleoside inhibitor of HIV reverse transcriptase, such as AZT, 3TC, ddC or ddI. A preferred inhibitor of HIV protease is indinavir, which is the sulfate salt of N-(2(R)-hydroxy-1-(S)-indanyl)-2(R)-phenylmethyl-4-(S)-hydroxy-5-(1-(4-(3-pyridyl-methyl)-2(S)-N'-(t-butylcarboxamido)-piperazinyl))-pentaneamide ethanolate, and is synthesized according to U.S. 5,413,999. Indinavir is generally administered at a dosage of 800 mg three times a day. Other preferred protease inhibitors are nelfinavir and ritonavir. Another preferred inhibitor of HIV protease is saquinavir which is administered in a dosage of 600 or 1200 mg tid. Preferred non-nucleoside inhibitors of HIV reverse transcriptase include efavirenz. The preparation of ddC, ddI and AZT are also described in EPO 0,484,071. These combinations may have unexpected effects on limiting the spread and degree of infection of HIV. Preferred combinations include those with the following (1) indinavir with efavirenz, and, optionally, AZT and/or 3TC and/or ddI and/or ddC; (2) indinavir, and any of AZT and/or ddI and/or ddC and/or 3TC, in particular, indinavir and AZT and 3TC; (3) stavudine and 3TC and/or zidovudine; (4) zidovudine and lamivudine and 141W94 and 1592U89; (5) zidovudine and lamivudine.

The pharmaceutical compositions of this invention may be administered orally, parenterally or via an implanted reservoir. Oral administration or administration by injection are preferred. In some cases, the pH of the formulation may be adjusted with pharmaceutically acceptable acids, bases or buffers to enhance the stability of the formulated compound or its delivery form. The term parenteral as

used herein includes subcutaneous, intracutaneous, intravenous, intramuscular, intra-articular, intrasynovial, intrasternal, intrathecal, and intralesional injection or infusion techniques.

5 When orally administered, the pharmaceutical compositions of this invention may be administered in any orally acceptable dosage form including, but not limited to, capsules, tablets, and aqueous suspensions and solutions. In the case of tablets for oral use, carriers which are commonly used include lactose and corn starch. Lubricating agents, such as magnesium stearate, are also typically added. For oral
10 administration in a capsule form, useful diluents include lactose and dried corn starch. When aqueous suspensions are administered orally, the active ingredient is combined with emulsifying and suspending agents. If desired, certain sweetening and/or flavoring and/or coloring agents may be added. Other suitable carriers for the above noted compositions can be found in standard pharmaceutical texts, e.g. in
15 "Remington's Pharmaceutical Sciences", 19th ed., Mack Publishing Company, Easton, Penn., 1995.

 The pharmaceutical compositions can be prepared by known procedures using well-known and readily available ingredients. The compositions of this invention
20 may be formulated so as to provide quick, sustained or delayed release of the active ingredient after administration to the patient by employing procedures well known in the art. In making the compositions of the present invention, the active ingredient will usually be admixed with a carrier, or diluted by a carrier, or enclosed within a carrier which may be in the form of a capsule, sachet, paper or other container.
25 When the carrier serves as a diluent, it may be a solid, semi-solid or liquid material which acts as a vehicle, excipient or medium for the active ingredient. Thus, the compositions can be in the form of tablets, pills, powders, beadlets, lozenges, sachets, elixirs, suspensions, emulsions, solutions, syrups, aerosols, (as a solid or in a liquid medium), soft and hard gelatin capsules, suppositories, sterile injectable solutions,
30 sterile packaged powders and the like. Further details concerning the design and preparation of suitable delivery forms of the pharmaceutical compositions of the invention are known to those skilled in the art.

Dosage levels of between about 0.01 and about 1000 milligram per kilogram ("mg/kg") body weight per day, preferably between about 0.5 and about 250 mg/kg body weight per day of the compounds of the invention are typical in a monotherapy
5 for the prevention and treatment of HIV mediated disease. Typically, the pharmaceutical compositions of this invention will be administered from about 1 to about 5 times per day or alternatively, as a continuous infusion. Such administration can be used as a chronic or acute therapy. The amount of active ingredient that may be combined with the carrier materials to produce a single dosage form will vary
10 depending upon the host treated and the particular mode of administration.

As the skilled artisan will appreciate, lower or higher doses than those recited above may be required. Specific dosage and treatment regimens for any particular patient will depend upon a variety of factors, including the activity of the specific
15 compound employed, the age, body weight, general health status, sex, diet, time of administration, rate of excretion, drug combination, the severity and course of the infection, the patient's disposition to the infection and the judgment of the treating physician. Generally, treatment is initiated with small dosages substantially less than the optimum dose of the peptide. Thereafter, the dosage is increased by small
20 increments until the optimum effect under the circumstances is reached. In general, the compound is most desirably administered at a concentration level that will generally afford antivirally effective results without causing any harmful or deleterious side effects.

25 When the compositions of this invention comprise a combination of a compound of the invention and one or more additional therapeutic or prophylactic agent, both the compound and the additional agent are usually present at dosage levels of between about 10 to 100%, and more preferably between about 10 and 80% of the dosage normally administered in a monotherapy regimen.

30

When these compounds or their pharmaceutically acceptable enantiomers, diastereomers, salts, solvates or prodrugs are formulated together with a

pharmaceutically acceptable carrier, the resulting composition may be administered in vivo to mammals, such as man, to inhibit HIV integrase or to treat or prevent HIV virus infection.

5 Accordingly, another aspect of this invention provides methods of inhibiting HIV integrase activity in patients by administering a compound of the present invention or a pharmaceutically acceptable enantiomer, diastereomer, salt or solvate thereof.

10 In one aspect of the invention, there is provided a method of treating an HIV infection in a patient, comprising administering to the patient a therapeutically effective amount of the compound of the invention, or a pharmaceutically acceptable enantiomer, diastereomer, solvate, prodrug or salt thereof.

15 Preferably, the method of administering the compound is effective to inhibit the function of the HIV integrase. In a preferred aspect, the method further comprises administering another compound having anti-HIV activity (as described above) prior to, after or concurrently with a compound of the invention.

20 The compounds of the invention may also be used as laboratory reagents. Compounds may be instrumental in providing research tools for designing of viral replication assays, validation of animal assay systems and structural biology studies to further enhance knowledge of the HIV disease mechanisms. Further, the compounds of the present invention are useful in establishing or determining the
25 binding site of other antiviral compounds, for example, by competitive inhibition.

 The compounds of this invention may also be used to treat or prevent viral contamination of materials and therefore reduce the risk of viral infection of laboratory or medical personnel or patients who come in contact with such materials,
30 e.g., blood, tissue, surgical instruments and garments, laboratory instruments and garments, and blood collection or transfusion apparatuses and materials.

Further, the compounds and compositions of the invention can be used for the manufacture of a medicament for treating HIV infection in a patient.

5

EXAMPLES

The specific examples that follow illustrate the syntheses of the compounds of the instant invention, and are not to be construed as limiting the invention in sphere or scope. The methods may be adapted to variations in order to produce compounds embraced by this invention but not specifically disclosed. Further, variations of the methods to produce the same compounds in somewhat different manner will also be evident to one skilled in the art.

In the following experimental procedures, all temperatures are understood to be in Centigrade (C) when not specified. The nuclear magnetic resonance (NMR) spectral characteristics refer to chemical shifts (δ) expressed in parts per million (ppm) versus tetramethylsilane (TMS) as reference standard. The relative area reported for the various shifts in the proton NMR spectral data corresponds to the number of hydrogen atoms of a particular functional type in the molecule. The nature of the shifts as to multiplicity is reported as broad singlet (bs or br s), broad doublet (bd or br d), broad triplet (bt or br t), broad quartet (bq or br q), singlet (s), multiplet (m), doublet (d), quartet (q), triplet (t), doublet of doublet (dd), doublet of triplet (dt), and doublet of quartet (dq). The solvents employed for taking NMR spectra are acetone- d_6 (deuterated acetone), DMSO- d_6 (perdeuterodimethylsulfoxide), D_2O (deuterated water), $CDCl_3$ (deuteriochloroform) and other conventional deuterated solvents.

The abbreviations used herein are conventional abbreviations widely employed in the art. Some of which are: calcd (calculated); DMSO (dimethylsulfoxide); EtOAc (ethyl acetate); HPLC (high-pressure liquid chromatography); LC/MS (liquid chromatography, mass spectroscopy); LDA (lithium diisopropyl amide); LiHMDS (lithium bis(trimethylsilyl)amide); SiO_2 (silica gel); THF (tetrahydrofuran), TFA (trifluoroacetic acid), Me (methyl), Et (ethyl), Ph (phenyl), tBuOK (potassium tert-

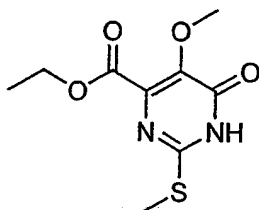
butoxide), NaOMe (sodium methoxide), NaOEt (sodium ethoxide), Boc (tert-butoxycarbonyl), and DEAD (diethylazo dicarboxylate).

5

EXAMPLE 1

Preparation of N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide

10 Compound 1A: 5-Methoxy-2-methylsulfanyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester

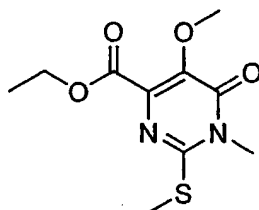


A solution of diethyl oxalate (27.2 g, 0.186 mol) and methyl methoxyacetate
15 (19.38 g, 0.186 mol) in dry tetrahydrofuran (250 ml) was treated at 22 °C with
sodium hydride (8.15 g of a 60 % dispersion in mineral oil, 0.203 mol) and the
resulting mixture was stirred at 22 °C for 18h. The tetrahydrofuran was then
evaporated under reduced pressure and the residual orange syrup was dissolved in a
solution of sodium ethoxide (0.189 mol, prepared from 4.35 g of sodium) in ethanol
20 (300 ml). Then powdered 2-methyl-2-thiopseudourea sulfate (25.9 g, 0.093 mol) was
added and the resulting mixture was stirred at 22 °C for 1 h and then heated at 60 °C
for 2 h. Acetic acid (5 ml) was then added to the gel like reaction mixture and the
ethanol was evaporated under reduced pressure. The residual paste was partitioned
between water and dichloromethane and the aqueous phase was extracted five times
25 with dichloromethane. The combined organic phases were washed with brine and
dried over anhydrous magnesium sulfate. Evaporation of the solvent gave a semi-
solid residue which was triturated with ethyl acetate (100 ml). The light yellow solid
formed was collected (10.02 g) and chromatographed on silica gel. Elution with a
mixture of dichloromethane and ethanol (98:2) gave 6.02 g (13 % yield) of the title

ester as white needles; mp 171–173 °C (ethyl acetate). ¹HNMR 400 MHz (CDCl₃) δ (ppm) : 1.42 (3H, t, J = 7.1 Hz, CH₃), 2.62 (3H, s, SCH₃), 3.99 (3H, s, OCH₃), 4.43 (2H, q, J = 7.1 Hz, CH₂), 11.8 (1H, broad s, NH). Anal. Calcd for C₉H₁₂N₂O₄S: C 44.25, H 4.95, N 11.47. Found: C 44.22, H 5.04, N 11.46.

5

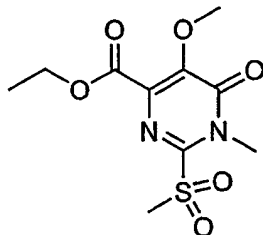
Compound 1B: 5-Methoxy-1-methyl-2-methylsulfanyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester



10

A solution of 5-methoxy-2-methylsulfanyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester (0.442 g, 1.81 mmol) in dry N,N-dimethylformamide (5 ml) was treated at 10 °C with sodium hydride (0.080 g of a 60 % dispersion in mineral oil, 2.0 mmol) and the resulting mixture was stirred for 10 min. Then
 15 iodomethane (0.123 ml, 2.0 mmol) was added and the mixture was stirred for another 1h. The solvent was then evaporated *in vacuo* and the residue was diluted with ethyl acetate, washed with saturated sodium bicarbonate, brine and dried over anhydrous magnesium sulfate. Evaporation of the solvent and chromatography of the residue on silica gel (elution toluene–ethyl acetate 9:1) gave 0.419 g (89 % yield) of the title
 20 ester as white needles; mp 74.5–76 °C (ether-hexane). ¹HNMR 400 MHz (CDCl₃) δ (ppm) : 1.41 (3H, t, J = 7.1 Hz, CH₃), 2.62 (3H, s, SCH₃), 3.55 (3H, s, NCH₃), 3.95 (3H, s, OCH₃), 4.43 (2H, q, J = 7.1 Hz, CH₂). Anal. Calcd for C₁₀H₁₄N₂O₄S: C 46.50, H 5.46, N 10.85. Found: C 46.73, H 5.21, N 10.93.

Compound 1C: 2-Methanesulfonyl-5-methoxy-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester

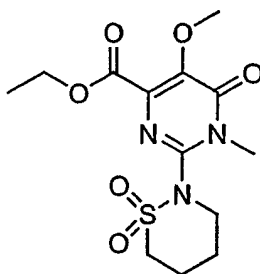


5

A solution of 5-methoxy-1-methyl-2-methylsulfonyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester (0.410 g, 1.58 mmol) in dry dichloromethane (30 ml) was treated at 20 °C with 3-chloroperoxybenzoic acid (0.71 g of 85 %, 3.5 mmol) and the resulting clear solution was stirred for 3 h. The reaction mixture was then diluted with ethyl acetate, washed successively with 10 % sodium bisulfite, saturated sodium bicarbonate, brine and dried over anhydrous magnesium sulfate. Evaporation of the solvent and chromatography of the residue on silica gel (elution toluene-ethyl acetate 9:1) gave 0.340 g (74 % yield) of the title ester as white needles; mp 75–76 °C (ethyl acetate-hexane). ¹HNMR 400 MHz (CDCl₃) δ (ppm) :
1.36 (3H, t, J = 7.1 Hz, CH₃), 3.43 (3H, s, SO₂CH₃), 3.84 (3H, s, NCH₃), 4.07 (3H, s, OCH₃), 4.37 (2H, q, J = 7.1 Hz, CH₂). Anal. Calcd for C₁₀H₁₄N₂O₆S: C 41.37, H 4.86, N 9.65. Found: C 41.21, H 4.64, N 9.55.

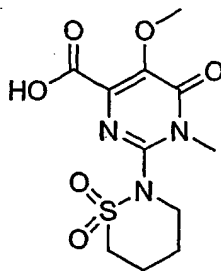
Compound 1D: 2-(1,1-Dioxo-1λ⁶-[1,2]thiazinan-2-yl)-5-methoxy-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester

20



A solution of 2-methanesulfonyl-5-methoxy-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester (0.180 g, 0.62 mmol) and [1,2]thiazinane 1,1-dioxide (0.100 g, 0.74 mmol) (W. Dirscherl and K. Otto, Chem. Ber. 89, 1956, 393) in dry N,N-dimethylformamide (2 ml) was cooled to 10 °C and treated under argon with sodium hydride (0.035 g of a 60 % dispersion in mineral oil, 0.9 mmol) and the resulting mixture was stirred for 1 h. The solvent was then evaporated *in vacuo*, the residue was dissolved in ethyl acetate, washed with saturated sodium bicarbonate, brine and dried over anhydrous magnesium sulfate. Evaporation of the solvent and chromatography of the residue on silica gel (elution toluene–ethyl acetate 7:3) gave 0.183 g (86 % yield) of the title ester as a clear oil. ¹HNMR 400 MHz (CDCl₃) δ (ppm) : 1.41 (3H, t, J = 7.1 Hz, CH₃), 1.65 (1H, m, CH), 2.3 – 2.6 (3H, m, CH₂ and CH), 3.2 (1H, m, CH), 3.6 (1H, m, CH), 3.64 (3H, s, NCH₃), 3.65 (1H, m, CH), 4.04 (3H, s, OCH₃), 4.05 (1H, m, CH), 4.41 (2H, q, J = 7.1 Hz, OCH₂). HRMS (FAB POS) calculated for C₁₃H₂₀N₃O₆S [M+H⁺] : 346.107282 : found : 346.107961.

Compound 1E: 2-(1,1-Dioxo-1λ⁶-[1,2]thiazinan-2-yl)-5-methoxy-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid



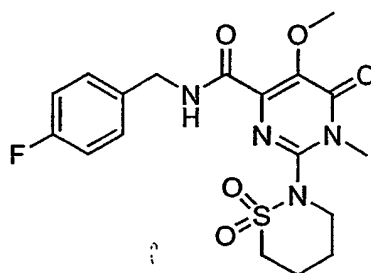
20

A solution of 2-(1,1-dioxo-1λ⁶-[1,2]thiazinan-2-yl)-5-methoxy-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester (0.180 g, 0.52 mmol) in 90 % ethanol (10 ml) was treated at 22 °C 1 N with sodium hydroxide (1 ml) and the resulting mixture was stirred for 30 min. Then 1 N hydrochloric acid (2 ml) was added and the ethanol was evaporated under reduced pressure. The residue was dissolved in ethyl acetate, washed with brine and dried over anhydrous magnesium

sulfate. Evaporation of the solvent and recrystallization of the solid residue from ethyl acetate gave 0.133 g (81 % yield) of the title acid as a white crystals; mp 180–181 °C. ¹HNMR 400 MHz (DMSO-d₆) δ (ppm) : 1.55–1.6 (1H, m, CH), 2.0–2.25 (3H, m, CH₂ and CH), 3.43 (2H, m, CH₂), 3.44 (3H, s, NCH₃), 3.75 (2H, m, CH₂), 3.83 (3H, s, OCH₃). Anal. Calcd for C₁₁H₁₅N₃O₆S: C 41.64, H 4.76, N 13.24. Found: C 41.84, H 4.96, N 12.88.

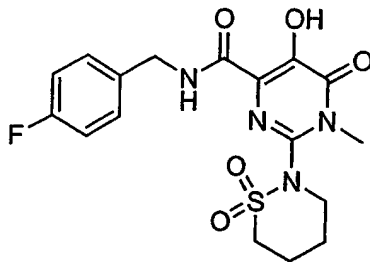
Compound 1F: 2-(1,1-Dioxo-1λ⁶-[1,2]thiazinan-2-yl)-5-methoxy-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid 4-fluorobenzylamide

10



A mixture of 2-(1,1-dioxo-1λ⁶-[1,2]thiazinan-2-yl)-5-methoxy-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid (0.235 g, 0.74 mmol) and 4-fluorobenzylamine (0.14 g, 1.13 mmol) in acetonitrile (10 ml) was treated at 22 °C with triethylamine (0.32 ml, 2.3 mmol) followed by benzotriazole-1-yloxy-tris-pyrrolidino-phosphonium hexafluorophosphate (0.42 g, 0.81 mmol) and the resulting mixture was stirred for 3 h. The reaction mixture was then diluted with ethyl acetate, washed with saturated sodium bicarbonate, brine and dried over anhydrous magnesium sulfate. Evaporation of the solvent and chromatography of the residue on silica gel (elution toluene–ethyl acetate 1:1 to ethyl acetate) gave 0.291 g (92 % yield) of the title amide as a white solid. ¹HNMR 400 MHz (CDCl₃) δ (ppm) : 1.6–1.7 and 2.3–2.5 (4H, m, 2 x CH₂), 3.2 and 3.5 (2 x 1H, m, CH₂), 3.62 (3H, s, NCH₃), 3.7 and 3.95 (2 x 1H, m, CH₂), 4.04 (3H, s, OCH₃), 4.59 (2H, d, J = 5.8 Hz, NCH₂), 7.04 (2H, m, aromatics), 7.31 (2H, m, aromatics), 7.75 (1H, broad, NH). HRMS (FAB POS) calculated for C₁₈H₂₂FN₄O₅S [M+H⁺] : 425.129495 : found : 425.129040.

Compound 1: N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide

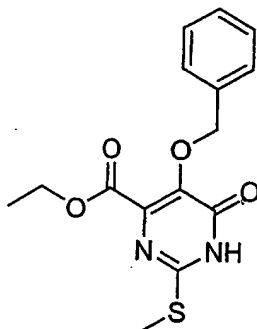


5

A mixture of 2-(1,1-dioxo-1 λ ⁶-[1,2]thiazinan-2-yl)-5-methoxy-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid 4-fluoro-benzylamide (0.115 g, 0.27 mmol) in 2,4,6-collidine (3 ml) was treated with lithium iodide (0.060 g) and the resulting mixture was heated at 120 °C for 30 min. The collidine was then evaporated
 10 *in vacuo*, the residue was diluted with a small volume of water and the pH was adjusted to a value of 7 with 0.1 N hydrochloric acid. The aqueous phase was extracted three times with dichloromethane and the combined organic extracts were washed with brine, dried over anhydrous magnesium sulfate and concentrated under reduced pressure. The residue was purified by chromatography on reversed phase
 15 silica gel (Waters, C-18, 125 Å, elution with water-acetonitrile 7:3 to 6:4) to give 0.058 g (52 % yield) of the title amide as a white solid; mp 226–227 °C. (dichloromethane-ethanol).

Compound 1G: 5-Benzyloxy-2-methylsulfanyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester

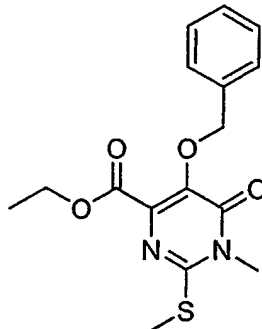
20



A solution of diethyl oxalate (21.06 g, 0.144 mol) and ethyl benzyloxyacetate (28.0 g, 0.144 mol) in dry tetrahydrofuran (200 ml) was treated at 22 °C with sodium hydride (6.34 g of a 60 % dispersion in mineral oil, 0.158 mol). Ethanol (0.05 ml) was added and the resulting mixture was stirred at 22 °C for 18h. The tetrahydrofuran was then evaporated under reduced pressure and the residual orange syrup was dissolved in a solution of sodium ethoxide (0.072 mol, prepared from 1.65 g of sodium) in ethanol (200 ml). Then powdered 2-methyl-2-thiopseudourea sulfate (20.1 g, 0.072 mol) was added and the resulting mixture was heated at 60 °C for 6 h. Acetic acid (5 ml) was then added to the gel like reaction mixture and the ethanol was evaporated under reduced pressure. The residual paste was partitioned between water and dichloromethane and the aqueous phase was extracted twice with dichloromethane. The combined organic phases were washed with saturated sodium bicarbonate, brine, dried over anhydrous magnesium sulfate and concentrated under reduced pressure. Chromatography on silica gel (elution with a gradient of ethyl acetate 0-20 % in toluene) followed by crystallization from ethyl acetate-hexane gave 8.34 g (18 % yield) of the title ester as white needles; mp 109–110 °C. ¹HNMR 400 MHz (CDCl₃) δ (ppm) : 1.35 (3H, t, J = 7.1 Hz, CH₃), 2.62 (3H, s, SCH₃), 4.37 (2H, q, J = 7.1 Hz, CH₂), 5.28 (2H, s, OCH₂), 7.35 – 7.52 (5H, m, aromatics), 12.2 (1H, broad s, NH). Anal. Calcd for C₁₅H₁₆N₂O₄S: C 56.23, H 5.03, N 8.74. Found: C 56.23, H 4.86, N 8.76.

Compound 1H: 5-Benzyloxy-1-methyl-2-methylsulfanyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester

25



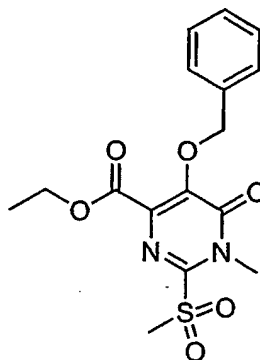
Method 1: 5-Benzyloxy-2-methylsulfanyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester (6.190 g, 19.32 mmol) in dry N,N-dimethylformamide (60 ml) was reacted at 10 °C with sodium hydride (0.90 g of a 60 % dispersion in mineral oil, 22.5 mmol) and iodomethane (1.4 ml, 21.7 mmol) and chromatographed to give first 0.581 g (9 % yield) of 5-benzyloxy-6-methoxy-2-methylsulfanyl-pyrimidine-4-carboxylic acid ethyl ester as white crystals; mp 48–49 °C (ethyl acetate-hexane). ¹HNMR 400 MHz (CDCl₃) δ (ppm) : 1.32 (3H, t, J = 7.1 Hz, CH₃), 2.56 (3H, s, SCH₃), 4.07 (3H, s, OCH₃), 4.36 (2H, q, J = 7.1 Hz, CH₂), 5.04 (2H, s, OCH₂), 7.3 – 7.45 (5H, m, aromatics). HRMS (FAB POS) calculated for C₁₆H₁₉N₂O₄S [M+H⁺] : 335.106554 : found : 335.106641, δ -0.3 ppm. The following fractions then gave 5.836 g (90 % yield) of the title ester as white needles; mp 84–85 °C (ethyl acetate-hexane). ¹HNMR 400 MHz (CDCl₃) δ (ppm) : 1.33 (3H, t, J = 7.0 Hz, CH₃), 2.62 (3H, s, SCH₃), 3.57 (3H, s, NCH₃), 4.35 (2H, q, J = 7.0 Hz, CH₂), 5.22 (2H, s, OCH₂), 7.3 – 7.5 (5H, m, aromatics). Anal. Calcd for C₁₆H₁₈N₂O₄S: C 57.47, H 5.43, N 8.38. Found: C 57.37, H 5.42, N 8.36.

Method 2: To a stirred solution of methylthiocyanate (91 g, 1.245 Moles) and N-methylhydroxylamine hydrochloride (100 g, 1.197 Moles) in 1:1 ethanol/water (v/v, 500 mL) was carefully added Na₂CO₃ (63.6 g, 0.6 Moles) over 30 min at room temperature. The resulting reaction mixture was stirred 48 h then cooled in an ice-water bath. To this was slowly added diethyl acetylenedicarboxylate (192 mL, 1.2 Moles) over 10 minutes. After stirring 2 h, the resulting dark-brown reaction mixture was transferred to a separatory funnel and extracted with ethyl acetate (3 X 250 mL). The combined organic layers were dried (Na₂SO₄/activated charcoal), filtered and concentrated to give dark-brown residue (310.1 g) which was used in the next step without further purification.

A solution of the above crude intermediate in xylenes (1.2 L) was placed in a pre-heated oil bath (140 °C) and stirred for 48 h. The mixture was then cooled and concentrated to give a dark solid. This crude product mixture was dissolved into

anhydrous DMF (1.5 L) to which was added benzyl bromide (119 mL, 1 Mole) followed by K_2CO_3 (138 g, 1 Mole). After stirring for 72 h at room temperature, the reaction mixture was diluted with hexanes (1 L) and filtered. The filtrate was concentrated under vacuum and the resulting residue was taken up into ether (1 L), washed with water (3 X 250 mL) and brine (200 mL). The organic layer was dried (Na_2SO_4 /activated charcoal), filtered and concentrated to afford dark paste which was dissolved in ether/hexanes (2:1, 1.5 L). Approximately 1L of the solvent was removed on a rotary evaporator. The resulting solution was left overnight at room temperature and the brown-crystals which formed were removed by filtration. The filtrate was concentrated then chromatographed (EM Science silica gel 60, column size $\sim 30 \times 10 \text{ cm}^2$) using 8:1:1 Hexanes/ETOAc/ CH_2Cl_2 solvent system and collected into 250 mL fractions. The fractions containing the desired compound were combined and concentrated to give a viscous oil which was dissolved into Hex/ Et_2O ($\sim 4:1$) and allowed to crystallize (92.3 g). An additional 8.84 grams of product was isolated after the mother liquor was concentrated and allowed to sit for 48h. Resulting in a total yield 101.14 g (25.3% yield).

Compound 1I: 5-Benzyloxy-2-methanesulfonyl-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester

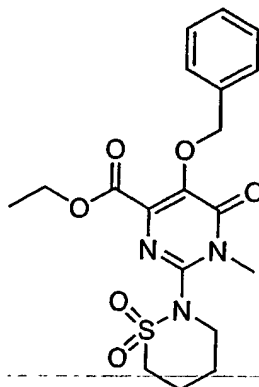


Reaction of 5-benzyloxy-1-methyl-2-methylsulfonyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester (5.83 g, 17.43 mmol) in dry dichloromethane (250 mL) with 3-chloroperoxybenzoic acid (12.5 g of 85 %, 61.6 mmol) gave 5.27 g (82 % yield) of the title ester as white needles; mp 65 – 67 °C

(ethyl acetate-hexane). ^1H NMR 400 MHz (CDCl_3) δ (ppm) : 1.30 (3H, t, $J = 7.1$ Hz, CH_3), 3.45 (3H, s, SO_2CH_3), 3.87 (3H, s, NCH_3), 4.32 (2H, q, $J = 7.1$ Hz, CH_2), 5.42 (2H, s, OCH_2), 7.3 – 7.5 (5H, m, aromatics). Anal. Calcd for $\text{C}_{16}\text{H}_{18}\text{N}_2\text{O}_6\text{S}$: C 52.45, H 4.95, N 7.65. Found: C 52.27, H 4.69, N 7.58.

5

Compound 1J: 2-(1,1-Dioxo-1 λ ⁶-[1,2]thiazinan-2-yl)-5-benzyloxy-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester



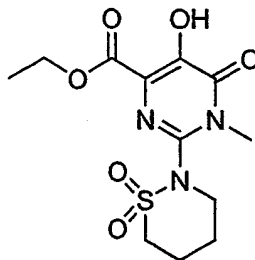
10

Reaction of 2-methanesulfonyl-5-benzyloxy-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester (2.21 g, 6.03 mmol) and [1,2]thiazinane 1,1-dioxide (0.97 g, 7.2 mmol) in dry N,N-dimethylformamide (30 ml) with sodium hydride (0.35 g of a 60 % dispersion in mineral oil, 8.8 mmol) gave 2.35 g (92 % yield) of the title ester as white needles; mp 109–110 °C. ^1H NMR 400 MHz (CDCl_3) δ (ppm) : 1.29 (3H, t, $J = 7.1$ Hz, CH_3), 1.62 (1H, m, CH), 2.3 – 2.6 (3H, m, CH_2 and CH), 3.15 – 3.21 (1H, m, CH), 3.5 – 3.6 (1H, m, CH), 3.64 (3H, s, NCH_3), 3.64 (1H, m overlapping with NCH_3 , CH), 3.97–4.05 (1H, m, CH), 4.32 (2H, q, $J = 7.1$ Hz, OCH_2), 5.29 (2H, AB system, $J_{\text{AB}} = 10.6$ Hz, $\Delta\nu = 62.5$ Hz, OCH_2), 7.3 – 7.5 (5H, m, aromatics). Anal. Calcd for $\text{C}_{19}\text{H}_{23}\text{N}_3\text{O}_6\text{S}$: C 54.15, H 5.50, N 9.97. Found: C 53.88, H 5.40, N 9.91.

15

20

Compound 1K: 2-(1,1-Dioxo-1 λ ⁶-[1,2]thiazinan-2-yl)-5-hydroxy-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester



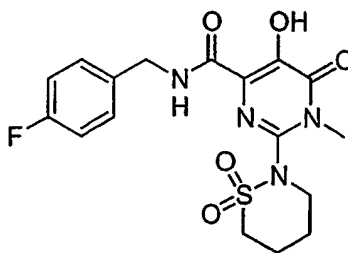
5

A solution of 2-(1,1-dioxo-1 λ ⁶-[1,2]thiazinan-2-yl)-5-benzyloxy-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester (4.05 g, 9.61 mmol) in a mixture of ethanol (250 ml) and ethyl acetate (250 ml) was hydrogenated over 10 % palladium on activated carbon (0.9 g) and under 1 atmosphere of hydrogen for 1 h.

- 10 The catalyst was then filtered and the solvent was evaporated under reduced pressure. Recrystallization of the residue from ethyl acetate gave 2.63 g (82 % yield) of the title ester as white crystals; mp 186–189 °C. ¹HNMR 400 MHz (CDCl₃) δ (ppm) : 1.42 (3H, t, J = 7.0 Hz, CH₃), 1.6–1.63 (1H, m, CH), 2.3–2.6 (3H, m, CH₂ and CH), 3.2 (1H, m, CH), 3.55 – 3.75 (2H, m, CH₂), 3.65 (3H, s, NCH₃), 3.95 – 4.05 (1H, m, CH), 4.44 (2H, q, J = 7.0 Hz, OCH₂). HRMS (FAB POS), calculated for C₁₂H₁₈N₃O₆S [M+H⁺] : 332.091632 : found: 332.092342.
- 15

Compound 1: N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide

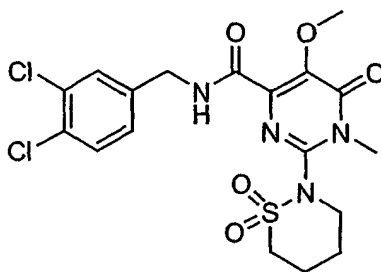
20



A mixture of 2-(1,1-dioxo-1 λ^6 -[1,2]thiazinan-2-yl)-5-hydroxy-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester (1.00 g, 3.02 mmol) and 4-fluorobenzylamine (1.31 g, 10.5 mmol) in a mixture of anhydrous ethyl alcohol (40 ml) and N,N-dimethylformamide (10 ml) was heated under reflux for 10 h. The solvent was then evaporated *in vacuo* and the residue was diluted with dichloromethane (500 ml). The organic phase was washed with 5 % aqueous acetic acid, brine and dried over anhydrous sodium sulfate. Evaporation of the solvent and recrystallization of the white solid obtained from dichloromethane and ethanol gave 1.04 g (84 % yield) of the title amide as white crystals; mp 227-228 °C. ¹HNMR 400 MHz (CDCl₃) δ (ppm): 1.75 and 2.07 (2x1H, m, CH₂), 2.28-2.43 (2H, m, CH₂), 3.19-3.31 (2H, m, CH₂), 3.64 (3H, s, NCH₃), 3.6-3.8 (2H, m, CH₂), 4.59 (2H, AB part of ABX system, J_{AB} = 14.9 Hz, Δv = 36.1 Hz, J_{AX} = 6.6 Hz, J_{BX} = 5.8 Hz, NCH₂), 7.06 (2H, m, aromatics), 7.31 (2H, m, aromatics), 7.52 (1H, broad NH). Anal. Calcd for C₁₇H₁₉FN₄O₅S: C 49.75, H 4.67, N 13.65. Found: C 49.67, H 4.63, N 13.77.

EXAMPLE 2

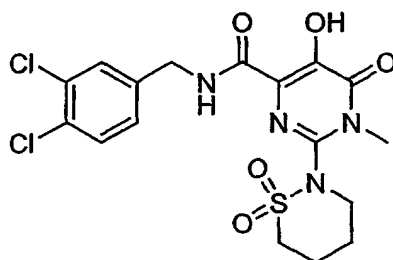
Compound 2A: 2-(1,1-Dioxo-1 λ^6 -[1,2]thiazinan-2-yl)-5-methoxy-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid 3,4-dichloro-benzylamide



Reaction of 2-(1,1-dioxo-1 λ^6 -[1,2]thiazinan-2-yl)-5-methoxy-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid (0.130 g, 0.41 mmol) and 3,4-dichlorobenzylamine (0.105 g, 0.60 mmol) in acetonitrile (10 ml) with triethylamine (0.16 ml, 1.14 mmol) and benzotriazole-1-yloxy-tris-pyrrolidino-phosphonium hexafluorophosphate (0.23 g, 0.44 mmol) 0.195 g (100 % yield) of the title amide as

a white solid; mp 212 °C.(dec.). ¹HNMR 400 MHz (DMSO-d₆) δ (ppm) : 1.55 – 1.6 and 2.1 – 2.25 (1H and 3H, 2 m, 2 x CH₂), 3.4-3.5 (2H, m, CH₂), 3.46 (3H, s, NCH₃), 3.75-3.85 (2H, m, CH₂), 3.84 (3H, s, OCH₃), 4.46 (2H, d, J = 6.0 Hz, NCH₂), 7.32 (1H, dd, J = 2.0 Hz and J = 8.3 Hz, aromatic), 7.56 (1H, d, J = 2.0 Hz, aromatic), 7.63 (1H, d, J = 8.3 Hz, aromatic), 9.03 (1H, broad t, NH). HRMS (FAB POS) calculated for C₁₈H₂₁Cl₂N₄O₅S [M+H⁺] : 475.060972 : found : 475.060157, δ 1.7 ppm.

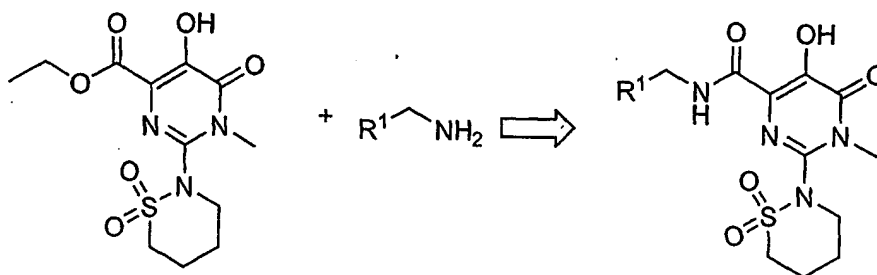
Compound 2: N-[(3,4-dichlorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxo-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide



Reaction of 2-(1,1-dioxo-1λ⁶-[1,2]thiazinan-2-yl)-5-methoxy-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid 3,4-dichloro-benzylamide (0.160 g, 0.337 mmol) in 2,4,6-collidine with lithium iodide gave 0.118 g (76 % yield) of the title amide as a white solid. ¹HNMR 400 MHz (CDCl₃) δ (ppm) : 1.70 – 1.85 and 2.06 – 2.15 (2 x 1H, 2 m, CH₂), 2.3 – 2.4 (2H, m, CH₂), 3.2 – 3.33 (2H, m, CH₂), 3.64 (3H, s, NCH₃), 3.7-3.85 (2H, m, CH₂), 4.58 (2H, AB part of ABX system, J_{AB} = 15.5 Hz, Δv = 49.6 Hz, J_{AX} = 6.6 Hz, J_{BX} = 6.1 Hz, NCH₂), 7.18 (1H, broad dd, aromatic), 7.42 (1H, broad d, aromatic), 7.45 (1H, d, J = 8.1 Hz, aromatic, 7.57 (1H, broad NH). HRMS (FAB POS) calculated for C₁₇H₁₉Cl₂N₄O₅S [M+H⁺] : 461.045322 : found : 461.045599.

EXAMPLE 3

Preparation of compounds 3-49

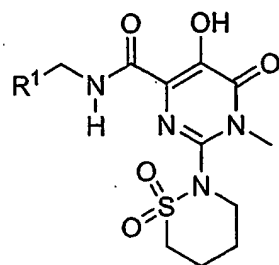


A solution of 2-(1,1-dioxo-1 λ^6 -[1,2]thiazinan-2-yl)-5-hydroxy-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester (0.045 mmol) and the
 5 selected amine (0.180 mmol) in a mixture of anhydrous ethyl alcohol (0.5 ml) and N,N-dimethylformamide (0.5 ml) was heated in a sealed vial at 85 °C for 8 h. Acetic acid (1 ml) was added and the products were purified on a Shimadzu automated preparative HPLC system (column YMC Pack C-18, 5 μ , 20 x 100 mm, elution water 0.1 % trifluoroacetic acid – acetonitrile). Spectrometry (MS) data were
 10 determined with a Micromass ZMD Platform TSQ 7000 LC/MS in positive electrospray mode.

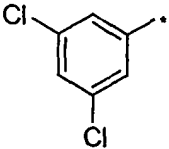
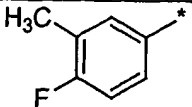
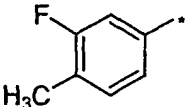
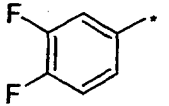
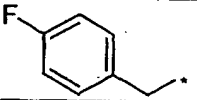
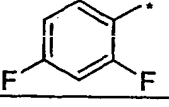
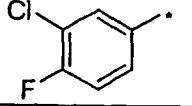
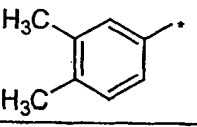
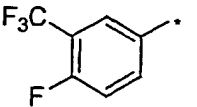
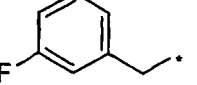
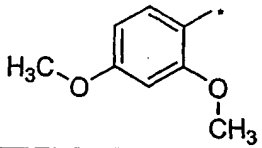
The collected compounds were analysed using the following LC/MS conditions.

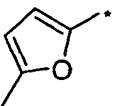
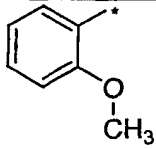
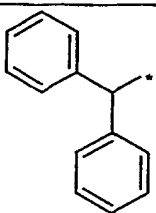
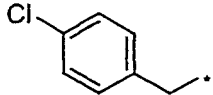
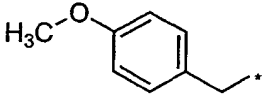
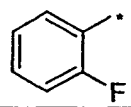
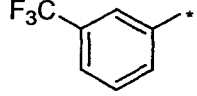
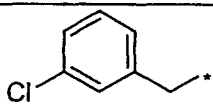
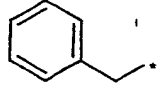
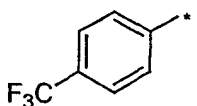
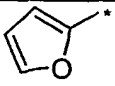
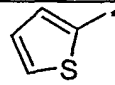
15

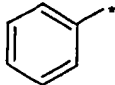
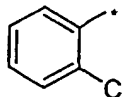
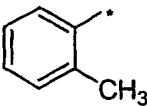
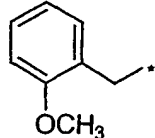
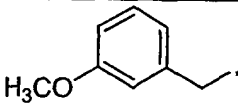
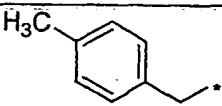
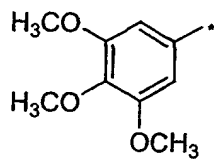
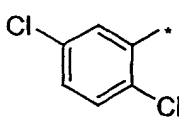
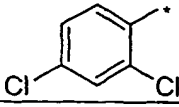
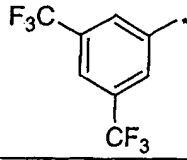
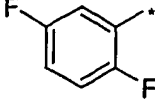
Column: Premisphere 5 μ C-8, 4.6 x 30 mm
 Solvent A: 10 % CH₃CN – 90 % H₂O , 0.05 % TFA
 Solvent B: 90 % CH₃CN – 10 % H₂O , 0.05 % TFA
 Gradient: 100 % A / 0 % B to 0 % A / 100 % B
 20 Gradient time: 2 minutes, hold time 1 minute.
 Flow rate: 4 ml/min.
 Detector wavelength: 220 nm.

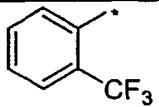
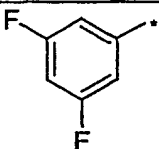
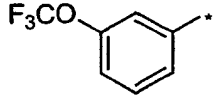
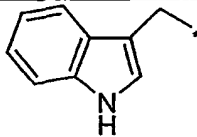


Compound	R ¹	HPLC Retention time (min)	Formula	MS Data (M+H) ⁺
3		1.71	C ₁₇ H ₁₉ FN ₄ O ₅ S	411
4		1.72	C ₁₇ H ₁₉ ClN ₄ O ₅ S	427
5		1.71	C ₁₇ H ₁₉ ClN ₄ O ₅ S	427
6		1.76	C ₁₉ H ₂₃ FN ₄ O ₅ S	439
7		1.93	C ₁₉ H ₂₂ Cl ₂ N ₄ O ₅ S	489
8		1.68	C ₁₈ H ₂₂ N ₄ O ₅ S	407
9		1.69	C ₁₈ H ₂₂ N ₄ O ₅ S	407
10		1.57	C ₁₈ H ₂₂ N ₄ O ₆ S	423
11		1.58	C ₁₈ H ₂₂ N ₄ O ₆ S	423

Compound	R ¹	HPLC Retention time (min)	Formula	MS Data (M+H) ⁺
12		1.84	C ₁₇ H ₁₈ Cl ₂ N ₄ O ₅ S	461
13		1.71	C ₁₈ H ₂₁ FN ₄ O ₅ S	425
14		1.72	C ₁₈ H ₂₁ FN ₄ O ₅ S	425
15		1.66	C ₁₇ H ₁₈ F ₂ N ₄ O ₅ S	429
16		1.67	C ₁₈ H ₂₁ FN ₄ O ₅ S	425
17		1.65	C ₁₇ H ₁₈ F ₂ N ₄ O ₅ S	429
18		1.74	C ₁₇ H ₁₈ ClFN ₄ O ₅ S	445
19		1.76	C ₁₉ H ₂₄ N ₄ O ₅ S	421
20		1.80	C ₁₈ H ₁₈ F ₄ N ₄ O ₅ S	479
21		1.74	C ₁₈ H ₂₁ FN ₄ O ₅ S	425
22		1.62	C ₁₉ H ₂₄ N ₄ O ₇ S	453

Compound	R ¹	HPLC Retention time (min)	Formula	MS Data (M+H) ⁺
23		1.55	C ₁₆ H ₂₀ N ₄ O ₆ S	397
24		1.63	C ₁₈ H ₂₂ N ₄ O ₆ S	423
25		1.89	C ₂₄ H ₂₆ N ₄ O ₅ S	483
26		1.77	C ₁₈ H ₂₁ ClN ₄ O ₅ S	441
27		1.62	C ₁₉ H ₂₄ N ₄ O ₆ S	437
28		1.61	C ₁₇ H ₁₉ FN ₄ O ₅ S	411
29		1.78	C ₁₈ H ₁₉ F ₃ N ₄ O ₅ S	461
30		1.76	C ₁₈ H ₂₁ ClN ₄ O ₅ S	441
31		1.65	C ₁₈ H ₂₂ N ₄ O ₅ S	407
32		1.80	C ₁₈ H ₁₉ F ₃ N ₄ O ₅ S	461
33		1.44	C ₁₅ H ₁₈ N ₄ O ₆ S	383
34		1.54	C ₁₅ H ₁₈ N ₄ O ₅ S ₂	399

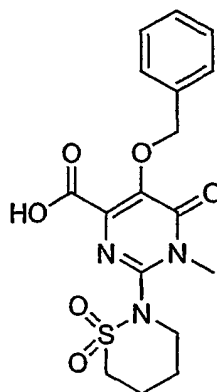
Compound	R ¹	HPLC Retention time (min)	Formula	MS Data (M+H) ⁺
35		1.58	C ₁₇ H ₂₀ N ₄ O ₅ S	393
36		1.70	C ₁₇ H ₁₉ ClN ₄ O ₅ S	427
37		1.66	C ₁₈ H ₂₂ N ₄ O ₅ S	407
38		1.68	C ₁₉ H ₂₄ N ₄ O ₆ S	437
39		1.64	C ₁₉ H ₂₄ N ₄ O ₆ S	437
40		1.74	C ₁₉ H ₂₄ N ₄ O ₅ S	421
41		1.47	C ₂₀ H ₂₆ N ₄ O ₈ S	483
42		1.80	C ₁₇ H ₁₈ Cl ₂ N ₄ O ₅ S	461
43		1.84	C ₁₇ H ₁₈ Cl ₂ N ₄ O ₅ S	461
44		1.95	C ₁₉ H ₁₈ F ₆ N ₄ O ₅ S	529
45		1.64	C ₁₇ H ₁₈ F ₂ N ₄ O ₅ S	429

Compound	R ¹	HPLC Retention time (min)	Formula	MS Data (M+H) ⁺
46		1.78	C ₁₈ H ₁₉ F ₃ N ₄ O ₅ S	461
47		1.67	C ₁₇ H ₁₈ F ₂ N ₄ O ₅ S	429
48		1.81	C ₁₈ H ₁₉ F ₃ N ₄ O ₆ S	477
49		1.64	C ₂₀ H ₂₃ N ₅ O ₅ S	446

EXAMPLE 4

Compound 50A: 5-Benzyloxy-2-(1,1-dioxo-1λ⁶-[1,2]thiazinan-2-yl)-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid

5

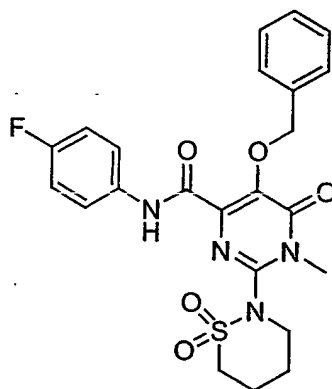


A solution of 5-benzyloxy-2-(1,1-dioxo-1λ⁶-[1,2]thiazinan-2-yl)-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester (1.02 g, 2.42 mmol) in 90
 10 % ethanol (30 ml) was treated with 1 N aqueous sodium hydroxide (5 ml) and stirred at 25 °C for 2 h. The ethanol was then evaporated *in vacuo*, the residue was acidified

with 1 N hydrochloric acid (8 ml) and extracted with ethyl acetate. The organic phase was washed with brine, dried over anhydrous sodium sulfate and concentrated.

Recrystallization of the solid residue from ethyl acetate gave 0.89 g (93 % yield) of the title ester as a white crystals; mp 161 – 163 °C. ¹HNMR 400 MHz (CDCl₃) δ (ppm) : 1.65 (1H, m, CH), 2.3 – 2.6 (3H, m, CH₂ and CH), 3.18 – 3.24 (1H, m, CH), 3.47 – 3.55 (1H, m, CH), 3.66 (3H, s, NCH₃), 3.6 – 3.7 (1H, m overlapping with NCH₃, CH), 3.96-4.03 (1H, m, CH), 5.56 (2H, AB system, J_{AB} = 10.8 Hz, Δv = 59.4 Hz, OCH₂), 7.35 – 7.5 (5H, m, aromatics). Anal. Calcd for C₁₇H₁₉N₃O₆S: C 51.90, H 4.86, N 10.68. Found: C 51.76, H 4.66, N 10.69.

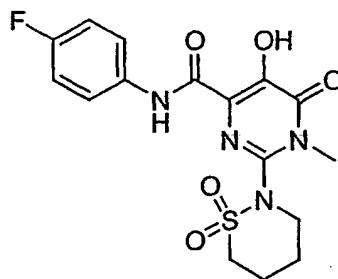
Compound 50B: 5-Benzyloxy-2-(1,1-dioxo-1λ⁶-[1,2]thiazinan-2-yl)-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid (4-fluoro-phenyl)-amide



A solution of 5-benzyloxy-2-(1,1-dioxo-1λ⁶-[1,2]thiazinan-2-yl)-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid (0.200 g, 0.51 mmol) in dichloromethane (20 ml) was treated at 25 °C with (chloromethylene)dimethylammonium chloride (0.080 g, 0.62 mmol) and the resulting mixture was stirred for 15 min. Then a solution of 4-fluoroaniline (0.117 g, 1.06 mmol) and pyridine (0.1 ml, 1.2 mmol) in dichloromethane (2 ml) was added and the reaction mixture was stirred for another 45 min. The mixture was quenched by addition of ethyl acetate and pH 6 phosphate buffer. The organic phase was then washed with saturated sodium bicarbonate, brine and dried over anhydrous magnesium sulfate. Evaporation of the solvent and chromatography of the residue on

silica gel (elution toluene – ethyl acetate 7:3) gave 0.270 g (88 % yield) of the title amide as white crystals; mp 201 – 202 °C.(ethyl acetate). ¹HNMR 400 MHz (CDCl₃) δ (ppm) : 1.65 – 1.69 and 2.35 – 2.7 (1H and 3H, 2 m, 2 x CH₂), 3.2-3.25 (1H, m, CH), 3.59 – 3.73 (2H, m, CH₂), 3.68 (3H, s, NCH₃), 3.99-4.07 (1H, m, CH), 5.47 (2H, AB system, J_{AB} = 10.8 Hz, Δv = 49.8 Hz, OCH₂), 6.93 (2H, m, aromatics), 7.22 (2H, m, aromatics), 7.39 – 7.52 (5H, m, aromatics) Anal. Calcd for C₂₃H₂₃FN₄O₅S: C 56.78, H 4.77, N 11.52. Found: C 56.83, H 4.58, N 11.53.

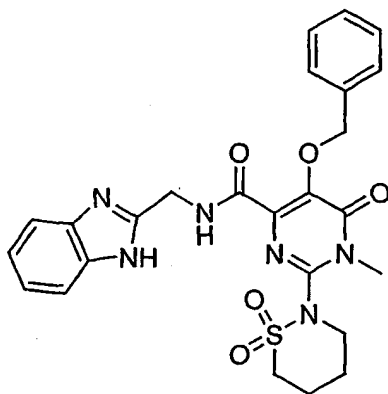
Compound 50: 2-(1,1-Dioxo-1λ⁶-[1,2]thiazinan-2-yl)-5-hydroxy-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid (4-fluoro-phenyl)-amide



Hydrogenolysis of 5-benzyloxy-2-(1,1-dioxo-1λ⁶-[1,2]thiazinan-2-yl)-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid (4-fluoro-phenyl)-amide (0.177 g, 0.364 mmol) gave 0.111 g (77 % yield) of the title amide as a white solid; mp 285 – 291 °C (dec.). ¹HNMR 400 MHz (CDCl₃) δ (ppm) : 1.83 – 1.92 and 2.14 – 2.25 (2 x 1H, 2 m, CH₂), 2.4 – 2.5 (2H, m, CH₂), 3.26 – 3.4 (2H, m, CH₂), 3.67 (3H, s, NCH₃), 3.7-3.9 (2H, m, CH₂), 7.11 (2H, m, aromatics), 7.58 (2H, m, aromatics), 9.0 (1H, broad s, NH). HRMS (FAB POS) calculated for C₁₆H₁₈FN₄O₅S [M+H⁺] : 397.098195 : found : 397.097570.

EXAMPLE 4

Compound 51A: 5-Benzyloxy-2-(1,1-dioxo-1 λ^6 -[1,2]thiazinan-2-yl)-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid (1H-benzimidazol-2-ylmethyl)-amide

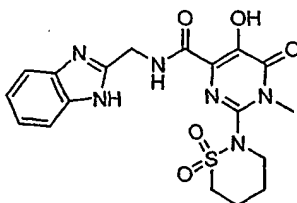


5

Coupling of 5-benzyloxy-2-(1,1-dioxo-1 λ^6 -[1,2]thiazinan-2-yl)-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid (0.250 g, 0.63 mmol) with 2-(aminomethyl)benzimidazole dihydrochloride hydrate (0.30 g, 1.26 mmol) gave 10 0.090 g (27 % yield) of the title amide as a white solid; mp 189 – 191 °C. ¹HNMR 400 MHz (CDCl₃) δ (ppm) : 1.67 and 2.35 – 2.52 (1H and 3H, 2 m, 2 x CH₂), 3.2-3.25 (1H, m, CH), 3.5 – 3.55 (1H, m, CH), 3.64 (3H, s, NCH₃), 3.69 (1H, m, CH), 3.94 (1H, m, CH), 4.67 (1H, d, J = 6.0 Hz, NCH₂), 5.38 (2H, AB system, J_{AB} = 11 Hz, $\Delta\nu$ = 48 Hz; OCH₂), 7.1 – 7.6 (9H, m, aromatics). HRMS (FAB POS), calculated for C₂₅H₂₇N₆O₅S [M+H⁺] : 523.176365 : found : 523.176120, δ 0.5 ppm.

15

Compound 51: N-(1H-benzimidazol-2-ylmethyl)-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide



20

Hydrogenolysis of 5-benzyloxy-2-(1,1-dioxo-1 λ^6 -[1,2]thiazinan-2-yl)-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid (1H-benzoimidazol-2-ylmethyl)-amide (0.078 g, 0.149 mmol) gave 0.030 g (46 % yield) of the title amide as a white solid. ¹HNMR 400 MHz (DMSO-d₆) δ (ppm) : 1.56 and 2.15-2.19 (1H and 3H, 2 m, 2 x CH₂), 3.41 (1H, m, CH), 3.45 (3H, s, NCH₃), 3.68 (1H, m, CH), 3.75 (2H, m, CH₂), 4.75 (2H, m, NCH₂), 7.15 (2H, m, aromatics), 7.51 (2H, m, aromatics). HRMS (FAB POS) calculated for C₁₈H₂₁N₆O₅S [M+H⁺] : 413.129415 : found : 413.129071.

10

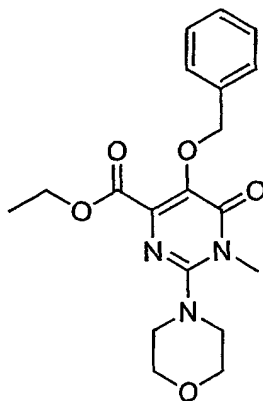
EXAMPLE 5

Preparation of compounds 52A-57

A representative procedure is shown for compound 52

Compound 52A: 5-Benzyloxy-1-methyl-2-morpholin-4-yl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester

15

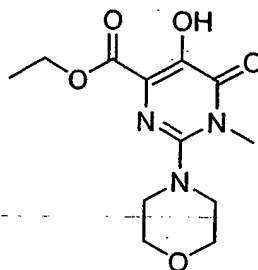


5-Benzyloxy-2-methanesulfonyl-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester (0.215 g, 0.58 mmol) was reacted with morpholine as described in the preparation of compound 1J. Chromatography on silica gel (elution toluene – ethyl acetate 8:2 to 6:4) gave first 0.050 g (30 % yield) of 5-benzyloxy-1-methyl-2,6-dioxo-1,2,3,6-tetrahydro-pyrimidine-4-carboxylic acid ethyl ester as white crystals; mp 184 – 187 °C (ethyl acetate). ¹HNMR 400 MHz (CDCl₃) δ (ppm) : 1.34 (3H, t, J = 7.1 Hz, CH₃), 3.40 (3H, s, NCH₃), 4.39 (2H, q, J = 7.1 Hz, OCH₂), 5.16 (3H, s, OCH₂), 7.35 – 7.5 (5H, m, aromatics), 8.12 (1H, broad s, NH). Anal.

Calcd for $C_{15}H_{16}N_2O_5$: C 59.20, H 5.30, N 9.20. Found: C 59.20, H 5.08, N 9.19. The end fractions then gave 0.102g (47% yield) of the title ester as a white syrup. ^1H NMR 400 MHz (CDCl_3) δ (ppm) : 1.29 (3H, t, $J = 7.1$ Hz, CH_3), 3.20 (4H, m, NCH_2), 3.53 (3H, s, NCH_3), 3.82 (4H, m, OCH_2), 4.32 (2H, q, $J = 7.1$ Hz, OCH_2), 5.17 (2H, s, OCH_2), 7.29 – 7.49 (5H, m, aromatics). HRMS (FAB POS) calculated for $C_{19}H_{24}N_3O_5$ [$\text{M} + \text{H}^+$] : 374.171596 : found : 374.1709485

Compound 52B: 5-Hydroxy-1-methyl-2-morpholin-4-yl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester

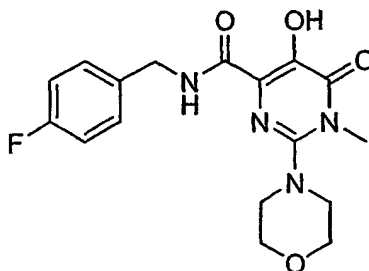
10



Hydrogenolysis of 5-benzyloxy-1-methyl-2-morpholin-4-yl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester (0.64 g, 1.71 mmol) gave 0.437 g (90 % yield) of the title ester as white crystals; mp 175 – 176 °C (ethanol). ^1H NMR 400 MHz (CDCl_3) δ (ppm) : 1.43 (3H, t, $J = 7.1$ Hz, CH_3), 3.13 (4H, m, NCH_2), 3.56 (3H, s, NCH_3), 3.83 (4H, m, OCH_2), 4.44 (2H, q, $J = 7.1$ Hz, OCH_2), 10.32 (1H, s, OH). Anal. Calcd for $C_{12}H_{17}N_3O_5$: C 50.88, H 6.04, N 14.83. Found: C 50.79, H 5.96, N 14.72.

20

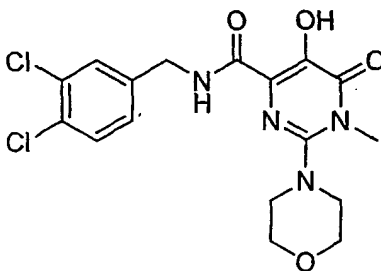
Compound 52: N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(4-morpholinyl)-6-oxo-4-pyrimidinecarboxamide



5

Reaction of 5-hydroxy-1-methyl-2-morpholin-4-yl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester (0.050 g, 0.176 mmol) with 4-fluorobenzylamine (0.07 g, 0.56 mmol) in ethyl alcohol (3 ml) gave 0.054 g (84 % yield) of the title ester as white crystals; mp 197 – 199 °C (ethyl acetate). ¹HNMR 400 MHz (CDCl₃) δ (ppm) : 3.08 (4H, m, NCH₂), 3.58 (3H, s, NCH₃), 3.85 (4H, m, OCH₂), 4.61 (2H, d, J = 6.0 Hz, NCH₂), 7.08 (2H, m, aromatics), 7.34 (2H, m, aromatics), 7.77 (1H, broad t, NH), 11.74 (1H, s, OH). Anal. Calcd for C₁₇H₁₉FN₄O₄: C 56.34, H 5.28, N 15.46. Found: C 56.39, H 5.02, N 15.45.

15 Compound 53: N-[(3,4-dichlorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(4-morpholinyl)-6-oxo-4-pyrimidinecarboxamide

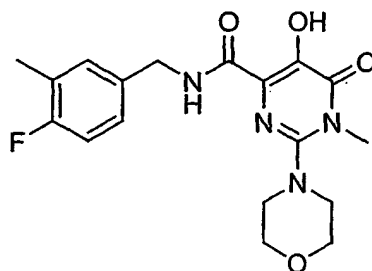


20

White crystals; mp 100 – 102 °C (ethyl acetate-hexane). ¹HNMR 400 MHz (CDCl₃) δ (ppm) : 3.07 (4H, m, NCH₂), 3.56 (3H, s, NCH₃), 3.83 (4H, m, OCH₂), 4.57 (2H, d, J = 6.5 Hz, NCH₂), 7.18 (1H, dd, J = 2.0 Hz and J = 8.2 Hz, aromatic),

7.42 (1H, d, $J = 2.0$ Hz, aromatic), 7.44 (1H, d, $J = 8.2$ Hz, aromatic), 7.78 (1H, broad t, NH). HRMS (FAB POS) calculated for $C_{17}H_{19}Cl_2N_4O_4$ [$M+H^+$]: 413.078336 : found : 413.076964.

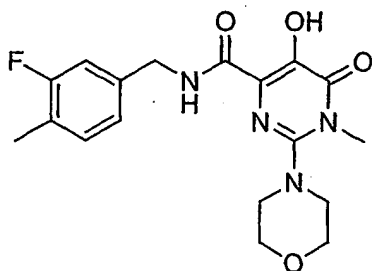
- 5 Compound 54: N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(4-morpholinyl)-6-oxo-4-pyrimidinecarboxamide



- 10 White crystals; mp 182 – 185 °C (ethyl acetate-hexane). ^1H NMR 400 MHz (CDCl_3) δ (ppm): 2.28 (3H, d, $J = 2.0$ Hz, CH_3), 3.05 (4H, m, NCH_2), 3.55 (3H, s, NCH_3), 3.82 (4H, m, OCH_2), 4.54 (2H, d, $J = 6.3$ Hz, NCH_2), 6.99 (1H, m, aromatic), 7.1 – 7.16 (2H, m, aromatics), 7.72 (1H, broad t, NH). Anal. Calcd for $C_{18}H_{21}FN_4O_4$: C 57.44, H 5.62, N 14.89. Found: C 57.31, H 5.46, N 14.79.

15

- Compound 55: N-[(3-fluoro-4-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(4-morpholinyl)-6-oxo-4-pyrimidinecarboxamide

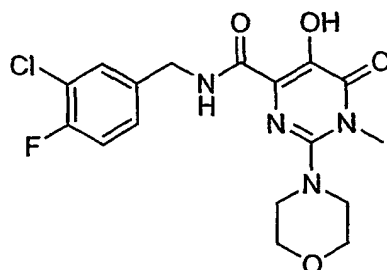


20

- White crystals; mp 166 – 167 °C (ethyl acetate-hexane). ^1H NMR 400 MHz (CDCl_3) δ (ppm): 2.27 (3H, d, $J = 1.8$ Hz, CH_3), 3.06 (4H, m, NCH_2), 3.56 (3H, s,

NCH₃), 3.83 (4H, m, OCH₂), 4.57 (2H, d, J = 6.3 Hz, NCH₂), 6.99 (2H, m, aromatics), 7.17 (1H, m, aromatic), 7.75 (1H, broad t, NH). Anal. Calcd for C₁₈H₂₁FN₄O₄: C 57.44, H 5.62, N 14.89. Found: C 57.29, H 5.62, N 14.82.

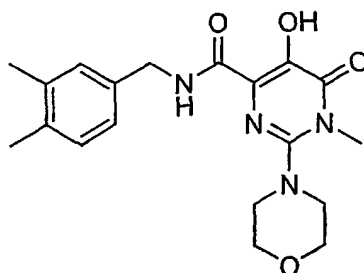
- 5 Compound 56: N-[(3-chloro-4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(4-morpholinyl)-6-oxo-4-pyrimidinecarboxamide



- 10 White crystals; mp 170 – 171 °C (ethyl acetate-hexane). ¹HNMR 400 MHz (CDCl₃) δ (ppm) : 3.07 (4H, m, NCH₂), 3.56 (3H, s, NCH₃), 3.83 (4H, m, OCH₂), 4.57 (2H, d, J = 6.2 Hz, NCH₂), 7.13 (1H, m, aromatic), 7.22 (1H, m, aromatic), 7.37 (1H, dd, J = 2.0 Hz and J = 7.0 Hz, aromatic), 7.78 (1H, broad t, NH). Anal. Calcd for C₁₇H₁₈ClFN₄O₄: C 51.45, H 4.57, N 14.12. Found: C 51.50, H 4.09, N 14.01. HRMS (FAB POS) calculated for C₁₇H₁₉ClFN₄O₄ [M+H⁺] : 397.107866 : found : 397.109213.
- 15

Compound 57: N-[(3,4-dimethylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(4-morpholinyl)-6-oxo-4-pyrimidinecarboxamide

20



White crystals; mp 164 – 166 °C (ethyl acetate-hexane). ¹HNMR 400 MHz (CDCl₃) δ (ppm) : 2.26 (3H, s, CH₃), 2.27 (3H, s, CH₃), 3.04 (4H, m, NCH₂), 3.55 (3H, s, NCH₃), 3.82 (4H, m, OCH₂), 4.55 (2H, d, J = 6.0 Hz, NCH₂), 7.06 – 7.24 (3H, m, aromatics), 7.72 (1H, broad t, NH). Anal. Calcd for C₁₉H₂₄N₄O₄: C 61.27, H 6.49, N 15.04. Found: C 61.07, H 6.83, N 14.92.

EXAMPLE 6

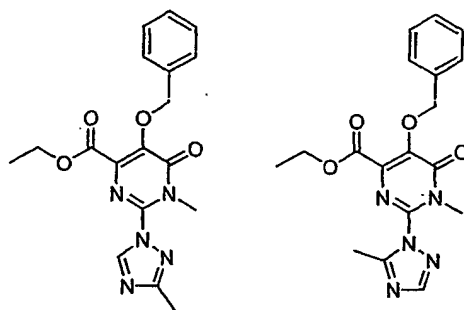
Preparation of compounds 58A-62

A representative procedure is shown for compound 58

10

Compounds 58A and 58B: 5-Benzyloxy-1-methyl-2-(3-methyl-[1,2,4]triazol-1-yl)-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester (58A) and 5-Benzyloxy-1-methyl-2-(5-methyl-[1,2,4]triazol-1-yl)-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester (58B)

15



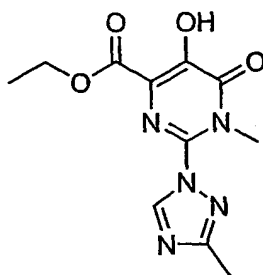
A solution of 5-benzyloxy-2-methanesulfonyl-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester (1.00 g, 2.73 mmol) and 3-methyl-1H-[1,2,4]triazole (0.272 g, 3.27 mmol) in dry N,N-dimethylformamide (10 ml) at 0 °C was treated with sodium hydride (0.16 g of a 60 % dispersion in mineral oil, 4.0 mmol) and the resulting mixture was stirred at 0 °C for 30 min and at 24 °C for another 30 min. The reaction mixture was then quenched with acetic acid (0.2 ml) and the solvent was evaporated *in vacuo*. The residue was dissolved in ethyl acetate, washed with saturated sodium bicarbonate, brine and dried over anhydrous magnesium sulfate. Evaporation of the solvent and chromatography on silica gel (elution toluene – acetone 90:10 to 85:15) first gave 0.358 g (35 % yield) of 5-benzyloxy-1-methyl-2-(3-methyl-[1,2,4]triazol-1-yl)-6-oxo-1,6-dihydro-pyrimidine-

20

25

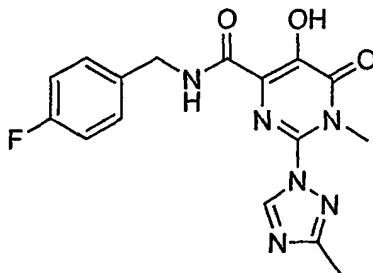
4-carboxylic acid ethyl ester as a clear oil. ^1H NMR 400 MHz (CDCl_3) δ (ppm) : 1.34 (3H, t, $J = 7.1$ Hz, CH_3), 2.51 (3H, s, CH_3), 3.73 (3H, s, NCH_3), 4.37 (2H, q, $J = 7.1$ Hz, CH_2), 5.37 (2H, s, OCH_2), 7.40 (3H, m, aromatics), 7.50 (2H, m, aromatics), 8.71 (1H, s, CH). HRMS (FAB POS) calculated for $\text{C}_{18}\text{H}_{20}\text{N}_5\text{O}_4$ [$\text{M}+\text{H}^+$] : 370.14 : found : 370.14. The following fractions then gave 0.395 g (39 % yield) of 5-benzyloxy-1-methyl-2-(5-methyl-[1,2,4]triazol-1-yl)-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester as a white solid. ^1H NMR 400 MHz (CDCl_3) δ (ppm) : 1.33 (3H, t, $J = 7.0$ Hz, CH_3), 2.67 (3H, s, CH_3), 3.52 (3H, s, NCH_3), 4.37 (2H, q, $J = 7.0$ Hz, CH_2), 5.41 (2H, s, OCH_2), 7.40 (3H, m, aromatics), 7.51 (2H, m, aromatics), 8.00 (1H, s, CH). HRMS (FAB POS) calculated for $\text{C}_{18}\text{H}_{20}\text{N}_5\text{O}_4$ [$\text{M}+\text{H}^+$] : 370.14 : found : 370.14.

Compound 58C: 5-Hydroxy-1-methyl-2-(3-methyl-[1,2,4]triazol-1-yl)-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester



Hydrogenolysis of 5-benzyloxy-1-methyl-2-(3-methyl-[1,2,4]triazol-1-yl)-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester (0.445 g, 1.20 mmol) in ethyl acetate) gave 0.286 g (85 % yield) of the title ester as white crystals; mp 160 – 162 °C. ^1H NMR 400 MHz (CDCl_3) δ (ppm) : 1.46 (3H, t, $J = 7.1$ Hz, CH_3), 2.51 (3H, s, CH_3), 3.68 (3H, s, NCH_3), 4.52 (2H, q, $J = 7.1$ Hz, CH_2), 8.63 (1H, s, CH), 10.91 (1H, s, OH). HRMS (FAB POS) calculated for $\text{C}_{11}\text{H}_{14}\text{N}_5\text{O}_4$ [$\text{M}+\text{H}^+$] : 280.10 : found : 280.10.

Compound 58: N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(3-methyl-1H-1,2,4-triazol-1-yl)-6-oxo-4-pyrimidinecarboxamide

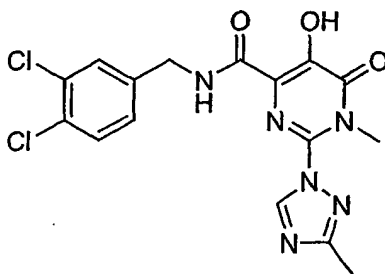


5

Reaction of 5-hydroxy-1-methyl-2-(3-methyl-[1,2,4]triazol-1-yl)-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester (0.060 g, 0.21 mmol) with 4-fluorobenzylamine (0.16 g, 1.28 mmol) gave 0.037 g (49 % yield) of the title ester as white crystals. ¹HNMR 400 MHz (CDCl₃) δ (ppm) : 2.51 (3H, s, CH₃), 3.63 (3H, s, NCH₃), 4.60 (2H, d, J = 6.2 Hz, NCH₂), 7.07 (2H, m, aromatics), 7.33 (2H, m, aromatics), 7.60 (1H, broad t, NH), 8.50 (1H, s, CH), 12.45 (1H, s, OH). HRMS (FAB POS) calculated for C₁₆H₁₆FN₆O₃ [M+H⁺] : 359.12 : found : 359.12.

Compound 59: N-[(3,4-dichlorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(3-methyl-1H-1,2,4-triazol-1-yl)-6-oxo-4-pyrimidinecarboxamide

15

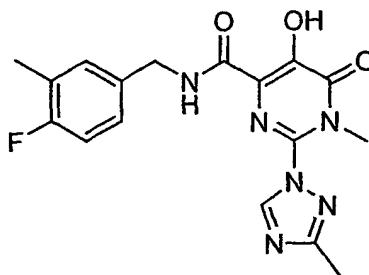


White crystals; mp 188 – 189 °C (dec.) (ethyl acetate-hexane). ¹HNMR 400 MHz (CDCl₃) δ (ppm) : 2.52 (3H, s, CH₃), 3.64 (3H, s, NCH₃), 4.59 (2H, d, J = 6.3 Hz, NCH₂), 7.20 (1H, dd, J = 2.0 Hz and J = 8.7 Hz, aromatic), 7.43 (1H, d, J = 2.0 Hz, aromatic), 7.46 (1H, dd, J = 8.7 Hz, aromatic), 7.65 (1H, broad t, NH), 8.52 (1H,

20

s, CH), 12.30 (1H, s, OH). HRMS (FAB POS) calculated for $C_{16}H_{15}Cl_2N_6O_3$ [$M+H^+$]
]: 409.05 : found : 409.05.

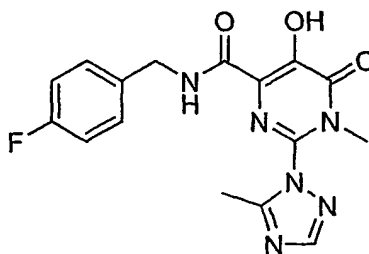
Compound 60: N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(3-methyl-1H-1,2,4-triazol-1-yl)-6-oxo-4-pyrimidinecarboxamide



White crystals; mp 204 – 205 °C (ethyl acetate-hexane). 1H NMR 400 MHz
($CDCl_3$) δ (ppm) : 2.29 (3H, d, $J = 1.9$ Hz, CH_3), 2.51 (3H, s, CH_3), 3.63 (3H, s,
10 NCH_3), 4.55 (2H, d, $J = 6.2$ Hz, NCH_2), 7.0 (1H, m, aromatic), 7.1 – 7.16 (2H, m,
aromatics), 7.57 (1H, broad t, NH), 8.50 (1H, s, CH), 12.48 (1H, s, OH). HRMS
(FAB POS) calculated for $C_{17}H_{18}FN_6O_3$ [$M+H^+$] : 373.13 : found : 373.13.

15

Compound 61: N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(5-methyl-1H-1,2,4-triazol-1-yl)-6-oxo-4-pyrimidinecarboxamide

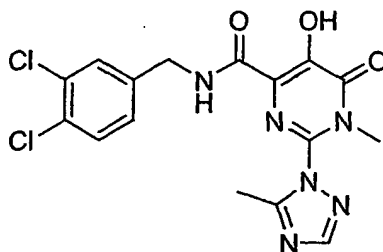


20

White crystals; mp 230 °C (ethyl acetate – ether). 1H NMR 400 MHz ($CDCl_3$)
 δ (ppm) : 2.54 (3H, s, CH_3), 3.42 (3H, s, NCH_3), 4.59 (2H, d, $J = 6.2$ Hz, NCH_2),

7.07 (2H, m, aromatics), 7.32 (2H, m, aromatics), 7.57 (1H, broad t, NH), 8.02 (1H, s, CH), 12.57 (1H, s, OH). HRMS (FAB POS), calculated for $C_{16}H_{16}FN_6O_3$ [$M+H^+$] : 359.126592 : found : 359.125641.

- 5 Compound 62: N-[(3,4-dichlorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(5-methyl-1H-1,2,4-triazol-1-yl)-6-oxo-4-pyrimidinecarboxamide

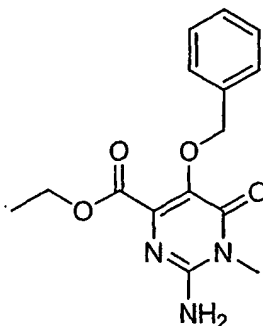


- 10 White crystals. 1H NMR 400 MHz ($CDCl_3$) δ (ppm) : 2.56 (3H, s, CH_3), 3.42 (3H, s, NCH_3), 4.58 (2H, d, $J = 6.2$ Hz, NCH_2), 7.18 (1H, dd, $J = 2.0$ Hz and $J = 8.1$ Hz, aromatic), 7.43 (1H, d, $J = 2.0$ Hz, aromatic), 7.45 (1H, dd, $J = 8.1$ Hz, aromatic), 7.64 (1H, broad t, NH), 8.03 (1H, s, CH), 12.42 (1H, s, OH). HRMS (FAB POS), calculated for $C_{16}H_{15}Cl_2N_6O_3$ [$M+H^+$] : 409.058269 : found : 409.056702.

15

EXAMPLE 7

Compound 63A: 2-Amino-5-benzyloxy-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester



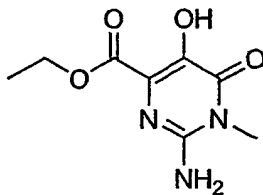
20

Gaseous ammonia was bubbled at 25 °C in a solution of 2-methanesulfonyl-5-benzyloxy-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester (0.200 g, 0.55 mmol) in acetonitrile (4 ml). After 30 min, the solvent was evaporated *in vacuo* and the residue was chromatographed on silica gel (elution toluene – ethyl acetate, 6:4 to 1:1) to give 0.099 g (59 % yield) of the title ester as a white solid.

¹HNMR 400 MHz (DMSO-d₆) δ (ppm) : 1.19 (3H, t, J = 7.1 Hz, CH₃), 3.29 (3H, s, NCH₃), 4.18 (2H, q, J = 7.1 Hz, OCH₂), 4.89 (2H, s, OCH₂), 7.21 (2H, broad s, NH₂), 7.31 – 7.37 (5H, m, aromatics). Anal. Calcd for C₁₅H₁₇N₃O₄: C 59.39, H 5.64, N 13.85. Found: C 59.31, H 5.46, N 13.72.

10

Compound 63B: 2-amino-5-hydroxy-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester.

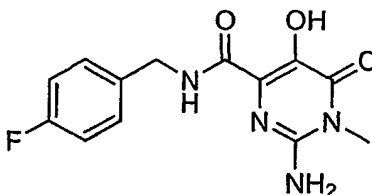


15

Hydrogenolysis of 2-amino-5-benzyloxy-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester (0.069 g, 0.23 mmol) gave 0.049 g of the title ester as an amorphous solid which was used as such for the next step. MS (ESI⁺) m/z 214 [M+H⁺].

20

Compound 63: 2-amino-5-hydroxy-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid 4-fluorobenzylamine

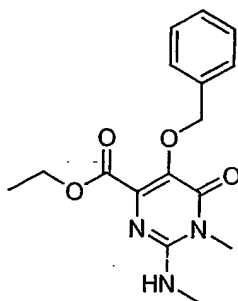


25

Reaction of 2-amino-5-hydroxy-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester (0.049 g, 0.23 mmol) with 4-fluorobenzylamine (0.10 g, 0.81 mmol) gave 0.030 g (42 % yield) of the title ester as a white solid. ¹HNMR 400 MHz (DMSO-d₆) δ (ppm) : 3.71 (3H, s, NCH₃), 4.45 (2H, d, J = 6.2 Hz, NCH₂), 6.51 (2H, broad s, NH₂), 7.1 – 7.19 (2H, m, aromatics), 7.34 – 7.38 (2H, m, aromatics), 8.71 (1H, broad t, NH). HRMS (FAB POS) calculated for C₁₃H₁₄FN₄O₃ [M+H⁺] : 293.104994 : found : 293.104843.

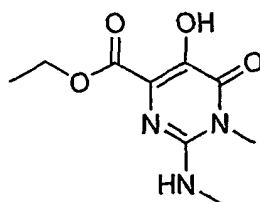
EXAMPLE 8

10 Compound 64A: 5-Benzyloxy-1-methyl-2-methylamino-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester



15 A solution of 2-methanesulfonyl-5-benzyloxy-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester (1.180 g, 3.22 mmol) in acetonitrile (20 ml) was treated at 25 °C with N,N-diisopropylethylamine (1.5 ml, 0.87 mmol) followed by 4.8 ml (9.6 mmol) of a 2 M solution of methylamine in tetrahydrofuran. After 30 min, the solvent was evaporated *in vacuo* and the residue was chromatographed on
20 silica gel (elution toluene – ethyl acetate, 25:75) to give 0.838 g (82 % yield) of the title ester as a clear syrup. ¹HNMR 400 MHz (CDCl₃) δ (ppm) : 1.30 (3H, t, J = 7.1 Hz, CH₃), 3.04 (3H, d, J = 5.1 Hz, NCH₃), 3.42 (3H, s, NCH₃), 4.32 (2H, q, J = 7.1 Hz, OCH₂), 4.58 (1H, broad q, NH), 5.07 (2H, s, OCH₂), 7.3 – 7.37 (3H, m, aromatics), 7.47 (2H, m, aromatics). HRMS (FAB POS) calculated for C₁₆H₂₀N₃O₄ [M+H⁺] : 318.145381 : found : 318.144860.
25

Compound 64B: 5-Hydroxy-1-methyl-2-methylamino-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester



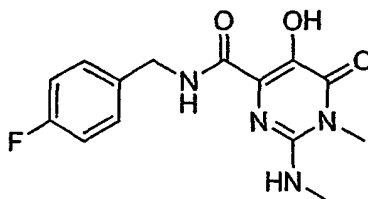
5

Hydrogenolysis of 5-benzyloxy-1-methyl-2-methylamino-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester (0.200 g, 0.63 mmol) gave 0.106 g (74 % yield) of the title ester as light yellow crystals; mp 226 °C (ethyl acetate). ¹HNMR 400 MHz (DMSO-d₆ + D₂O) δ (ppm) : 1.26 (3H, t, J = 7.1 Hz, CH₃), 2.75 (3H, s, NCH₃), 3.29 (3H, s, NCH₃), 4.26 (2H, q, J = 7.1 Hz, OCH₂). HRMS (FAB POS) calculated for C₉H₁₄N₃O₄ [M+H⁺] : 228.098431 : found : 228.099099.

10

Compound 64: 5-Hydroxy-1-methyl-2-methylamino-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid 4-fluoro-benzylamide

15



Reaction of 5-hydroxy-1-methyl-2-methylamino-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester (0.071 g, 0.231 mmol) with 4-fluorobenzylamine (0.14 g, 1.12 mmol) gave 0.080 g (84 % yield) of the title ester as a white solid. ¹HNMR 400 MHz (DMSO-d₆ + D₂O) δ (ppm) : 2.86 (3H, s, NCH₃), 3.39 (3H, s, NCH₃), 4.46 (2H, d, J = 6.1 Hz, NCH₂), 6.78 (1H exchanged D₂O, broad s, NH), 7.16 (2H, m, aromatics), 7.35 (2H, m, aromatics), 9.14 (1H, broad t, NH),

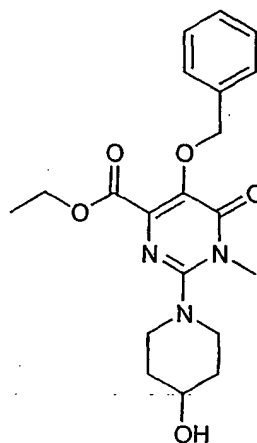
20

11.32 (1H exchanged D₂O, s, OH). HRMS (FAB POS) calculated for C₁₄H₁₆FN₄O₃ [M+H⁺]: 307.120644 : found : 307.119723.

EXAMPLE 9

5 Preparation of compounds 65A-66

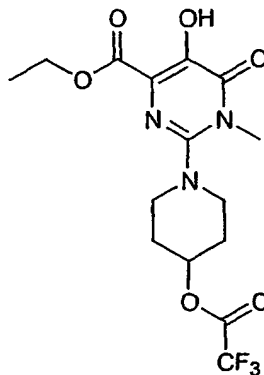
Compound 65A: 5-Benzyloxy-2-(4-hydroxypiperidin-1-yl)-1-methyl-6-oxo-1,6-dihydropyrimidine-4-carboxylic acid ethyl ester



10

A solution of 5-Benzyloxy-2-methanesulfonyl-1-methyl-6-oxo-1,6-dihydropyrimidine-4-carboxylic acid ethyl ester (0.725 g, 2 mmol) and 4-hydroxypiperidine (1 g, 10 mmol) in THF (20 mL) was stirred for 16 h at ambient temperature. Then, the reaction mixture was taken up in EtOAC (100 mL), washed with 1N aq HCl (2 X 15 10 mL) and brine (10 mL). The organic phase was dried (Na₂SO₄), filtered and concentrated to give the desired product as a pale yellow paste (0.608 g, 79%) which was used in the subsequent reaction without further purification. ¹H NMR (500 MHz, CDCl₃) δ: 7.46 (d, J = 7.0 Hz, 2H), 7.36-7.28 (m, 3H), 5.14 (s, 2H), 4.30 (q, J = 7.0 Hz, 2H), 3.92-3.87 (m, 1H), 3.50 (s, 3H), 3.44-3.39 (m, 2H), 3.01-2.95 (m, 2H), 20 2.79 (br s, 1H), 2.02-1.98 (m, 2H), 1.71-1.64 (m, 2H), 1.28 (t, J = 7.0 Hz, 3H). LRMS (M+H) calcd for C₂₀H₂₆N₃O₅: 388.44; found: 388.20

Compound 65B: 5-Hydroxy-1-methyl-6-oxo-2-[4-(2,2,2-trifluoroacetoxy)piperidin-1-yl]-1,6-dihydropyrimidine-4-carboxylic acid ethyl ester



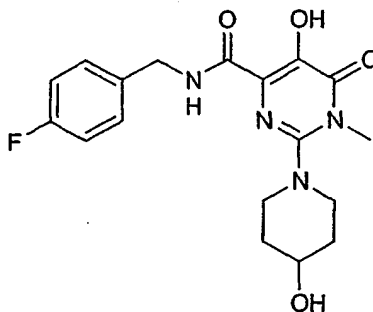
5

5-Benzyloxy-2-(4-hydroxypiperidin-1-yl)-1-methyl-6-oxo-1,6-dihydropyrimidine-4-carboxylic acid ethyl ester (0.760 g, 1.96 mmol) was treated with trifluoroacetic acid (20 mL) at ambient temperature. After 10 h, the reaction mixture was concentrated and purified on silica gel column using 2-5%

10 MeOH/CH₂Cl₂ to afford desired product as a brown paste (0.650 g, 84%). ¹H NMR (500 MHz, CDCl₃) δ: 10.02 (br s, 1H), 5.23-5.19 (m, 1H), 4.44 (q, J = 7.0 Hz, 2H), 3.57 (s, 3H), 3.36-3.31 (m, 2H), 3.15-3.10 (m, 2H), 2.17-2.11 (m, 2H), 2.02-1.95 (m, 2H), 1.42 (t, J = 7.0 Hz, 3H). LRMS (M+H) calcd for C₁₅H₁₉F₃N₃O₆: 394.32; found: 394.09

15

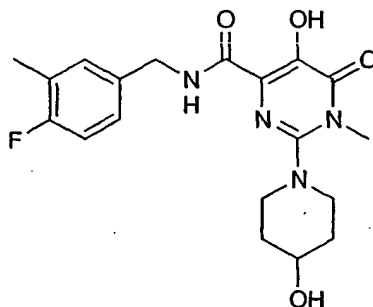
Compound 65: N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-2-(4-hydroxy-1-piperidinyl)-1-methyl-6-oxo-4-pyrimidinecarboxamide



20

A solution of 4-fluorobenzylamine (156 mg, 1.25 mmol) and 5-hydroxy-1-methyl-6-oxo-2-[4-(2,2,2-trifluoroacetoxy)piperidin-1-yl]-1,6-dihydropyrimidine-4-carboxylic acid ethyl ester (100 mg, 0.254 mmol) in dimethylformamide (3 mL) was heated at 120 °C for 2 h. Then, cooled and purified by preparative HPLC using
 5 MeOH/Water containing 0.1% TFA as an eluent. The fractions containing the product were combined and concentrated to afford title compound as a off-white solid (46 mg, 48%). ¹H NMR (500 MHz, CDCl₃) δ: 11.62 (s, 1H), 7.75 (br s, 1H), 7.31-7.28 (m, 2H), 7.04 (t, J = 8.7 Hz, 2H), 4.56 (d, J = 6.4 Hz, 2H), 3.95-3.89 (m, 1H), 3.52 (s, 3H), 3.28-3.23 (m, 2H), 2.88-2.83 (m, 2H), 2.02-1.96 (m, 2H), 1.73-1.66 (m, 2H),
 10 1.54 (br s, 1H). HRMS (M-H) calcd for C₁₈H₂₀FN₄O₄: 375.1469; found: 375.1472. Anal calcd for C₁₈H₂₁FN₄O₄: C, 57.44; H, 5.62; N, 14.88. Found: C, 57.19; H, 5.55; N, 14.82.

Compound 66: N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-2-(4-hydroxy-1-piperidinyl)-1-methyl-6-oxo-4-pyrimidinecarboxamide
 15



White solid. ¹H NMR (500 MHz, CDCl₃) δ: 11.65 (s, 1H), 7.74 (br s, 1H),
 20 7.14-7.09 (m, 2H), 6.97 (t, J = 8.9 Hz, 1H), 4.52 (d, J = 6.11 Hz, 1H), 3.95-3.89 (m, 1H), 3.52 (s, 3H), 3.28-3.22 (m, 2H), 2.89-2.84 (m, 2H), 2.27 (s, 3H), 2.02-1.96 (m, 2H), 1.73-1.66 (m, 2H), 1.52 (br s, 1H). HRMS (M-H) calcd for C₁₉H₂₂FN₄O₄: 389.1625; found: 389.1638. Anal calcd for C₁₉H₂₃FN₄O₄: C, 58.45; H, 5.93; N, 14.35. Found: C, 58.15; H, 5.84; N, 14.18.

25

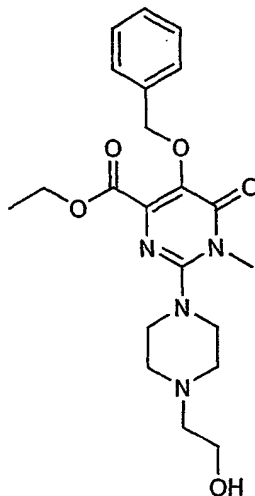
EXAMPLE 10

Preparation of compounds 67A-82

A representative procedure is shown for compound 67.

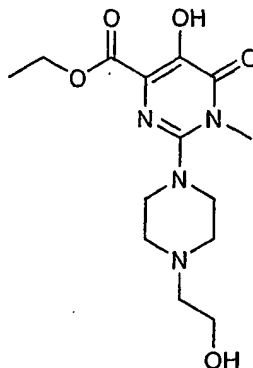
Compound 67A: 5-Benzyloxy-2-[4-(2-hydroxy-ethyl)-piperazin-1-yl]-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester

5



To a solution of 5-Benzyloxy-2-methanesulfonyl-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester (0.10 g, 0.27 mmol) dissolved in
10 THF (1 mL) was added 1-(2-hydroxyethyl)piperazine (0.05 mL, 0.41 mmol). The resulting mixture was stirred at 70° C for 3 hours. The mixture was concentrated and the resulting residue was purified by flash chromatography, eluting with 2% MeOH/CH₂Cl₂ to yield the title compound as a pale yellow oil (0.0423 g, 38% yield).
15 ¹H NMR (500 MHz, CDCl₃) δ: 7.45 (2H, d, *J*=7.0 Hz), 7.35-7.27 (3H, m), 5.14 (2H, s), 4.30 (2H, q, *J*=7.1 Hz), 3.63 (2H, t, *J*=5.3 Hz), 3.49 (3H, s), 3.21 (4H, t, *J*=4.7 Hz), 2.63 (4H, bs), 2.59 (2H, t, *J*=5.3 Hz), 1.27 (3H, t, *J*=7.2 Hz). HRMS (*M*+*H*) calcd for C₂₁H₂₉N₄O₅: 417.21381; found: 417.2147.

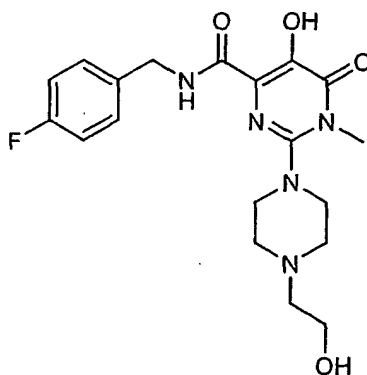
Compound 67B: 5-Hydroxy-2-[4-(2-hydroxy-ethyl)-piperazin-1-yl]-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester



5

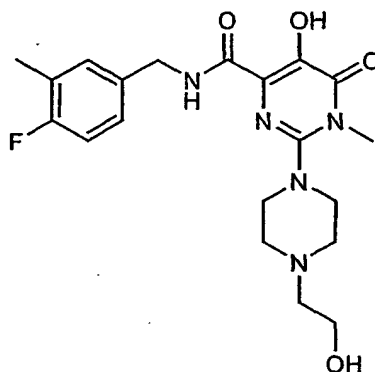
To a solution of 5-benzyloxy-2-[4-(2-hydroxy-ethyl)-piperazin-1-yl]-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester (0.2058 g, 0.49 mmol) dissolved in EtOAc (20 mL) was added a small amount of palladium on carbon. The mixture was shaken under H₂ at 40 psi for 10 hours; filtered over celite
 10 and concentrated to yield the title compound as a yellow waxy solid (0.0483 g, 30% yield). ¹H NMR (500 MHz, MeOD) δ: 4.44 (2H, q, *J*=7.1 Hz), 3.75 (2H, t, *J*=5.9 Hz), 3.56 (3H, s), 3.22-3.21 (4H, m), 2.78 (4H, bs), 2.68 (2H, t, *J*=5.9 Hz), 1.41 (3H, t, *J*=7.2 Hz). HRMS (*M*+*H*) calcd for C₁₄H₂₃N₄O₅: 327.16685; found: 327.1661.

15 Compound 67: N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-2-[4-(2-hydroxyethyl)-1-piperazinyl]-1-methyl-6-oxo-4-pyrimidinecarboxamide



5-Hydroxy-2-[4-(2-hydroxy-ethyl)-piperazin-1-yl]-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester (0.024 g, 0.074 mmol) was dissolved in DMF (1 mL) to which was added 4-fluoro-benzyl amine (0.047 mL, 0.37 mmol). The mixture was stirred at 70° C for 2 hours. The resulting mixture was purified by chromatography (YMC Combiprep ODS-A, 30mm x 50mm, MeOH/H₂O/0.1% TFA) to yield the title compound as a white powder (0.0149 g, 50% yield). ¹H NMR (500 MHz, MeOD) δ: 8.81 (1H, bs), 7.40 (2H, dd, *J*=8.5, 5.5 Hz), 7.08 (2H, t, *J*=8.8 Hz), 4.59 (2H, d, *J*=6.1 Hz), 3.94 (2H, t, *J*=5.2 Hz), 3.70 (4H, d, *J*=11.3 Hz), 3.58 (3H, s), 3.47-3.35 (6H, m). HRMS (M+H) calcd for C₁₉H₂₅FN₄O₅: 406.18907; found: 406.1895.

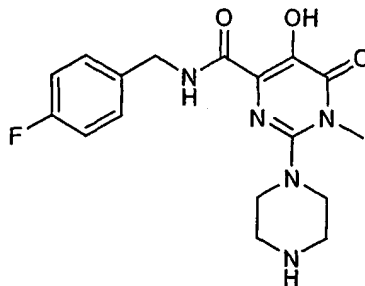
Compound 68: N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-2-[4-(2-hydroxyethyl)-1-piperazinyl]-1-methyl-6-oxo-4-pyrimidinecarboxamide



15

Colorless oil ¹H NMR (500 MHz, MeOD) δ: 8.79 (1H, bs), 7.24 (1H, d, *J*=7.3 Hz), 7.20-7.18 (1H, m), 6.99 (1H, t, *J*=9.0 Hz), 4.54 (2H, s), 3.94 (2H, s), 3.71-3.67 (4H, m), 3.58 (3H, s), 3.43-3.35 (6H, m), 2.26 (3H, d, *J*=1.5 Hz). HRMS (M+H) calcd for C₂₀H₂₇FN₄O₅: 420.20472; found: 420.2068.

Compound 69: N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(1-piperazinyl)-4-pyrimidinecarboxamide

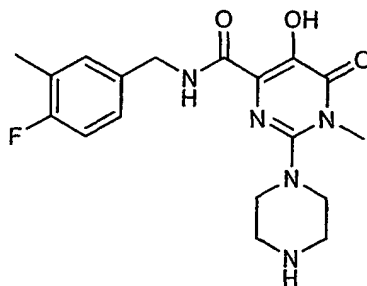


5

White solid, ^1H NMR (300 MHz, DMSO) δ : 9.28 (1H, t, $J=6.0$ Hz), 7.35 (2H, dd, $J=8.6, 5.7$ Hz), 7.16 (2H, $J=8.8$ Hz), 4.46 (2H, d, $J=6.2$ Hz), 3.39 (3H, s), 3.00-2.98 (4H, m), 2.84-2.81 (4H, m). HRMS ($M+H$) calcd for $\text{C}_{17}\text{H}_{22}\text{FN}_5\text{O}_3$: 362.16285; found: 362.1619. CHN theoretical: C, 56.50; H, 5.57; N, 19.38; F, 5.25; found: C, 56.29; H, 5.52; N, 19.18; F, 5.54.

10

Compound 70: N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(1-piperazinyl)-4-pyrimidinecarboxamide

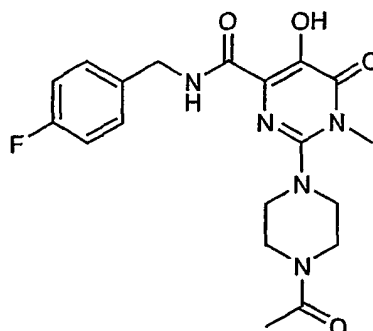


15

White solid ^1H NMR (500 MHz, DMSO) δ : 9.26 (1H, t, $J=5.0$ Hz), 7.22 (1H, dd, $J=7.6, 1.5$ Hz), 7.18-7.15 (1H, m), 7.08 (1H, t, $J=9.1$ Hz), 4.42 (2H, d, $J=6.4$ Hz), 3.39 (3H, s), 3.00-2.98 (4H, m), 2.84-2.82 (4H, m), 2.21 (3H, d, $J=1.5$ Hz). HRMS ($M+H$) calcd for $\text{C}_{18}\text{H}_{22}\text{FN}_5\text{O}_3$: 376.17850; found: 376.1783.

20

Compound 71: 2-(4-acetyl-1-piperazinyl)-N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-4-pyrimidinecarboxamide



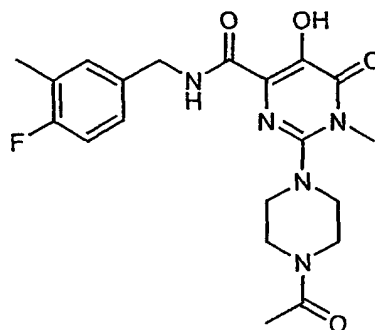
5

White solid ^1H NMR (500 MHz, CDCl_3) δ : 11.75 (1H, s), 7.68 (1H, s), 7.29 (2H, dd, $J=8.4, 5.3$ Hz), 7.04 (2H, t, $J=8.5$ Hz), 4.56 (2H, d, $J=6.1$ Hz), 3.74 (2H, s), 3.59 (2H, s), 3.55 (3H, s), 3.02 (4H, d, $J=17.7$ Hz), 2.11 (2H, s). HRMS ($\text{M}+\text{H}$) calcd for $\text{C}_{19}\text{H}_{23}\text{FN}_5\text{O}_4$: 404.17342; found: 404.1753. CHN theoretical 0.8 mol

10 H_2O : C, 54.62; H, 5.69; N, 16.76; F, 4.55; found: C, 54.62; H, 5.50; N, 16.41; F, 4.50.

Compound 72: 2-(4-acetyl-1-piperazinyl)-N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-4-pyrimidinecarboxamide

15

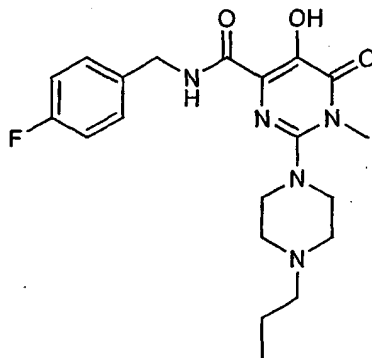


White solid ^1H NMR (500 MHz, CDCl_3) δ : 11.78 (1H, s), 7.66 (1H, s), 7.13 (1H, d, $J=7.0$ Hz), 7.11-7.08 (1H, m), 6.97 (1H, t, $J=8.8$ Hz), 4.52 (2H, d, $J=6.1$ Hz),

20 3.74 (2H, s), 3.60 (2H, s), 3.55 (3H, s), 3.02 (4H, d, $J=18.6$ Hz), 2.26 (3H, d, $J=1.8$ Hz), 2.12 (3H, s). HRMS ($\text{M}+\text{H}$) calcd for $\text{C}_{20}\text{H}_{25}\text{FN}_5\text{O}_4$: 418.18907; found:

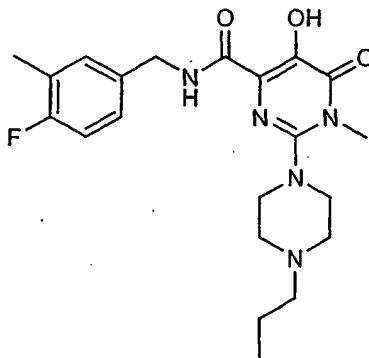
418.1885. CHN theoretical 0.5 mol H₂O: C, 56.33; H, 5.91; N, 16.42; F, 4.46;
found: C, 56.41; H, 5.68; N, 16.55; F, 4.64.

Compound 73: N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-
5 2-(4-propyl-1-piperazinyl)-4-pyrimidinecarboxamide



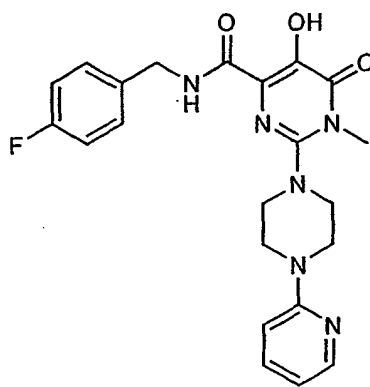
Brown oil ¹H NMR (500 MHz, CDCl₃) δ: 12.0 (1H, bs), 7.82 (1H, t, *J*=6.1
10 Hz), 7.29 (2H, dd, *J*=8.4, 5.3 Hz), 7.01 (3H, t, *J*=8.5 Hz), 4.54 (2H, d, *J*=6.4 Hz),
3.72 (2H, d, *J*=11.6 Hz), 3.49 (3H, s), 3.43-3.39 (4H, m), 3.04-2.97 (4H, m), 1.84-
1.76 (2H, m), 1.01 (3H, t, *J*=7.3 Hz). HRMS (*M*+*H*) calcd for C₂₀H₂₇FN₅O₃:
404.20980; found: 404.2095.

15 Compound 74: N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-
methyl-6-oxo-2-(4-propyl-1-piperazinyl)-4-pyrimidinecarboxamide



Brown oil ^1H NMR (500 MHz, CDCl_3) δ : 12.0 (1H, bs), 7.86 (1H, t, $J=5.8$ Hz), 7.12-7.07 (2H, m), 6.92 (1H, t, $J=8.8$ Hz), 4.48 (2H, d, $J=6.1$ Hz), 3.68 (2H, d, $J=10.7$ Hz), 3.46-3.40 (4H, m), 3.04-.01 (4H, m), 2.22 (3H, d, $J=1.5$ Hz), 1.81-1.74 (2H, m), 0.99 (3H, t, $J=7.3$ Hz). HRMS ($\text{M}+\text{H}$) calcd for $\text{C}_{21}\text{H}_{28}\text{FN}_5\text{O}_3$: 418.22545; found: 418.2247.

Compound 75: N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-[4-(2-pyridinyl)-1-piperazinyl]-4-pyrimidinecarboxamide

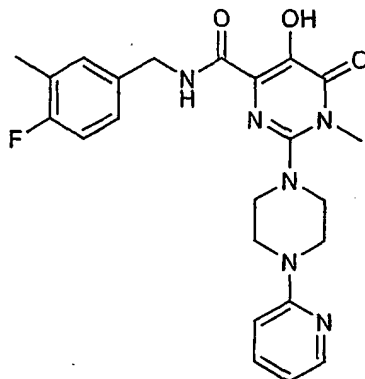


10

Pale pink foam ^1H NMR (500 MHz, CDCl_3) δ : 8.14 (1H, d, $J=5.8$ Hz), 7.93 (1H, t, $J=8.8$ Hz), 7.85 (1H, t, $J=6.2$ Hz), 7.31 (2H, dd, $J=8.5, 5.5$ Hz), 7.04-6.99 (3H, m), 6.95 (1H, t, $J=6.6$ Hz), 4.45 (2H, d, $J=6.4$ Hz), 3.90-3.88 (4H, m), 3.56 (3H, s), 3.30-3.28 (4H, m). HRMS ($\text{M}-\text{H}$) calcd for $\text{C}_{22}\text{H}_{22}\text{FN}_6\text{O}_3$: 437.17374; found: 437.1716.

15

Compound 76: N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-[4-(2-pyridinyl)-1-piperazinyl]-4-pyrimidinecarboxamide

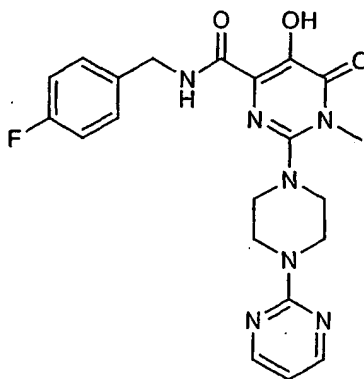


5

Pale pink foam ^1H NMR (500 MHz, CDCl_3) δ : 8.18 (1H, d, $J=5.8$ Hz), 7.93 (1H, t, $J=7.8$ Hz), 7.78 (1H, t, $J=6.1$ Hz), 7.15-7.10 (2H, m), 7.01 (1H, d, $J=9.1$ Hz), 6.97-6.94 (2H, m), 4.52 (2H, d, $J=6.4$ Hz), 3.91-3.89 (4H, m), 3.57 (3H, s), 3.30-3.28 (4H, m), 2.25 (3H, m). HRMS (M-H) calcd for $\text{C}_{23}\text{H}_{24}\text{FN}_6\text{O}_3$: 451.18939; found: 451.1894.

10

Compound 77: N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-[4-(2-pyrimidinyl)-1-piperazinyl]-4-pyrimidinecarboxamide

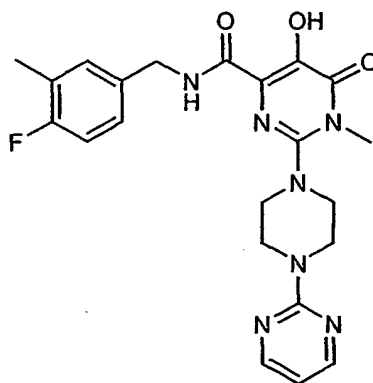


15

White solid ^1H NMR (300 MHz, CDCl_3) δ : 11.70 (bs), 8.33 (2H, d, $J=4.7$ Hz), 7.73 (1H, t, $J=6.2$ Hz), 7.29 (2H, dd, $J=8.6, 5.3$ Hz), 7.04 (2H, t, $J=8.6$ Hz), 6.56 (1H, t, $J=4.7$ Hz), 4.56 (2H, d, $J=6.2$ Hz), 3.97 (4H, t, $J=4.8$ Hz), 3.60 (3H, s), 3.10

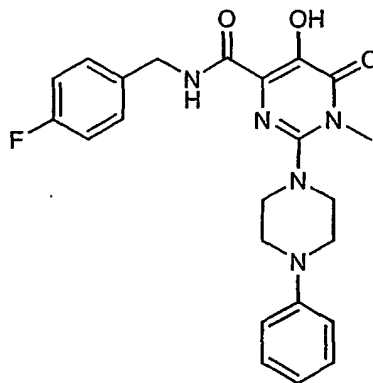
(4H, s, $J=5.1$ Hz). HRMS (M+H) calcd for $C_{21}H_{23}FN_7O_3$: 440.18465; found: 440.1847. CHN theoretical: C, 57.39; H, 5.04; N, 22.31; F, 4.32; found: C, 57.19; H, 5.03; N, 22.52; F, 4.42.

- 5 Compound 78: N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-[4-(2-pyrimidinyl)-1-piperazinyl]-4-pyrimidinecarboxamide



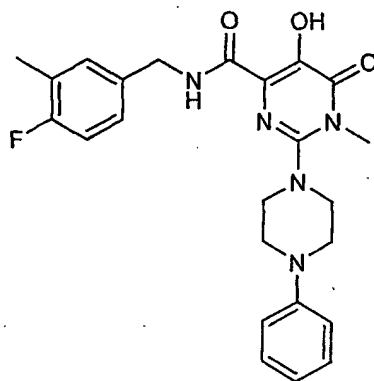
- 10 Lavender solid ^1H NMR (300 MHz, CDCl_3) δ : 11.92 (bs), 8.54 (2H, d, $J=4.8$ Hz), 7.72 (1H, t, $J=5.8$ Hz), 7.14-7.08 (2H, m), 6.95 (1H, t, $J=8.8$ Hz), 6.80 (1H, t, $J=3.6$ Hz), 4.51 (2H, d, $J=6.2$ Hz), 4.60 (4H, bs), 3.60 (3H, s), 3.20 (4H, bs), 2.24 (3H, s). HRMS (M+H) calcd for $C_{22}H_{25}FN_7O_3$: 454.20030; found: 454.1989.

- 15 Compound 79: N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(4-phenyl-1-piperazinyl)-4-pyrimidinecarboxamide



White solid ^1H NMR (300 MHz, CDCl_3) δ : 11.46 (1H, s), 7.77 (1H, t, $J=6.0$ Hz), 7.33-7.27 (4H, m), 7.05 (3H, t, $J=8.6$ Hz), 7.01-6.91 (3H, m), 4.58 (2H, d, $J=6.2$ Hz), 3.58 (3H, s), 3.33-3.30 (4H, m), 3.24-3.21 (4H, m). HRMS ($\text{M}+\text{H}$) calcd for $\text{C}_{23}\text{H}_{25}\text{FN}_5\text{O}_3$: 438.19415; found: 438.1955. CHN theoretical: C, 63.14; H, 5.52; N, 16.00; F, 4.34; found: C, 63.11; H, 5.47; N, 15.79; F, 4.36.

Compound 80: N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(4-phenyl-1-piperazinyl)-4-pyrimidinecarboxamide

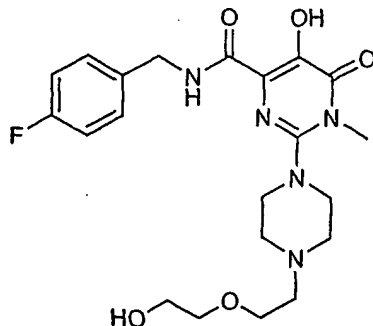


10

White solid ^1H NMR (300 MHz, CDCl_3) δ : 11.73 (1H, bs), 7.32-7.27 (3H, m), 7.15-7.05 (2H, m), 7.01-6.91 (3H, m), 4.54 (2H, d, $J=6.2$ Hz), 3.58 (3H, s), 3.33-3.31 (4H, m), 3.24-3.21 (4H, m), 2.26 (3H, s). HRMS ($\text{M}+\text{H}$) calcd for $\text{C}_{24}\text{H}_{27}\text{FN}_5\text{O}_3$: 452.20980; found: 452.2104.

15

Compound 81: N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-2-[4-[2-(2-hydroxyethoxy)ethyl]-1-piperazinyl]-1-methyl-6-oxo-4-pyrimidinecarboxamide

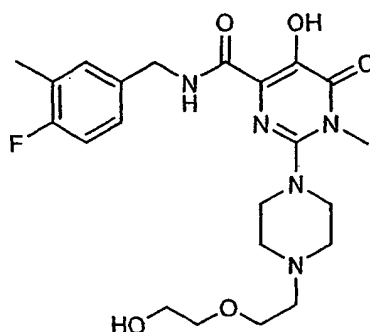


20

Colorless oil ^1H NMR (300 MHz, MeOD) δ : 8.83-8.79 (1H, m), 7.36 (2H, dd, $J=8.4, 5.5$ Hz), 7.04 (2H, t, $J=8.8$ Hz), 4.55 (2H, s), 3.90-3.84 (2H, m), 3.73-3.60 (6H, m), 3.55 (3H, s), 3.47-3.45 (2H, m), 3.40-3.34 (2H, m), 3.31-3.29 (4H, m).

5 HRMS (M-H) calcd for $\text{C}_{21}\text{H}_{27}\text{N}_5\text{O}_5\text{F}$: 448.19962; found: 448.2015.

Compound 82: N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-2-[4-[2-(2-hydroxyethoxy)ethyl]-1-piperazinyl]-1-methyl-6-oxo-4-pyrimidinecarboxamide



10

Colorless oil ^1H NMR (300 MHz, MeOD) δ : 8.77 (1H, bs), 8.12 (1H, s), 7.22-7.08 (2H, m), 6.95 (1H, t, $J=9.0$ Hz), 4.41 (2H, d, $J=51.2$ Hz), 3.91-3.84 (2H, m), 3.73-3.60 (6H, m), 3.54 (3H, s), 3.64 (2H, d, $J=4.4$ Hz), 3.39-3.23 (6H, m), 2.23 (3H, s). HRMS (M-H) calcd for $\text{C}_{22}\text{H}_{29}\text{N}_5\text{O}_5\text{F}$: 462.21527; found: 462.2147.

15

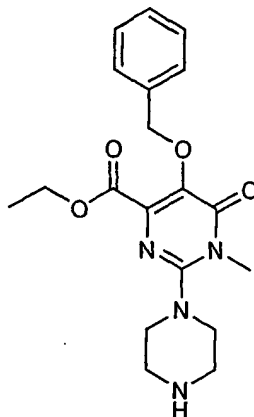
EXAMPLE 11

Preparation of compounds 83A-90

A representative procedure is given for compound 83

20

Compound 83A: 5-Benzyloxy-1-methyl-6-oxo-2-piperazin-1-yl-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester

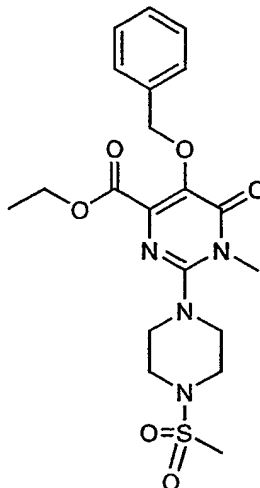


5

Following the procedure described for the synthesis of compound 67A 5-Benzyloxy-2-methanesulfonyl-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester (7.0 g, 19.2 mmol) was reacted with piperazine (17.0 g, 192 mmol) and to yield the title compound as a pale yellow oil (0.0418 g, 42% yield). ¹H NMR (500 MHz, CDCl₃) δ: 7.45 (2H, d, *J*=6.7 Hz), 7.34-7.27 (3H, m), 5.13 (2H, s), 4.30 (2H, q, *J*=7.2 Hz), 3.49 (3H, s), 3.15-3.13 (4H, m), 2.98-2.96 (4H, m), 1.89 (1H, bs), 1.27 (3H, *J*=7.2 Hz). HRMS (*M*+*H*) calcd for C₁₉H₂₅N₄O₄: 373.21414; found: 373.1866.

15

Compound 83B: 5-Benzyloxy-2-(4-methanesulfonyl-piperazin-1-yl)-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester

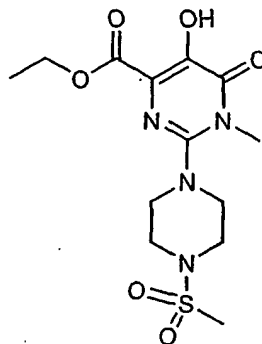


5

To a solution of ethyl 5-(benzyloxy)-1-methyl-6-oxo-2-(piperazin-1-yl)-1,6-dihydropyrimidine-4-carboxylate (0.145 g, 0.39 mmol) dissolved in pyridine (2 mL) was added methane sulfonyl chloride (0.31 mL, 0.39 mmol) and the resulting mixture was stirred at room temp for 2 hours. The mixture was diluted with EtOAc and the solids were removed by filtration and the solution was concentrated. The residue was purified by flash chromatography eluting with 0% to 50% to 100% EtOAc/hexane to give the title compound as a white foam (0.1364 g, 78% yield). ¹H NMR (300 MHz, CDCl₃) δ: 7.46 (2H, dd, *J*=7.9, 1.6 Hz), 7.37-7.27 (3H, m), 5.16 (2H, s), 4.30 (2H, q, *J*=7.2 Hz), 3.50 (3H, s), 3.34 (8H, d, *J*=6.2 Hz), 2.82 (3H, s), 1.28 (3H, t, *J*=7.1 Hz). HRMS (M+H) calcd for C₂₀H₂₇N₄O₆S: 451.16514; found: 451.1663.

10
15

Compound 83C: 5-Hydroxy-2-(4-methanesulfonyl-piperazin-1-yl)-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester



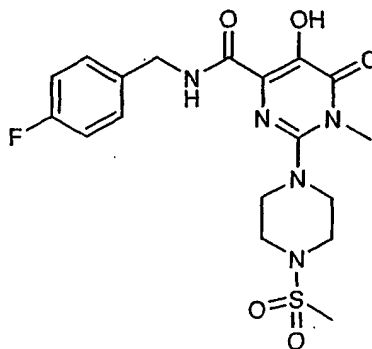
5

5-Benzyloxy-2-(4-methanesulfonyl-piperazin-1-yl)-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-carboxylic acid ethyl ester (0.22 g, 0.49 mmol) was treated with TFA to yield the title compound as a brown oily TFA salt (0.1564 g). ^1H NMR (500 MHz, CDCl_3) δ : 4.45 (2H, q, $J=7.1$ Hz), 3.59 (3H, s), 3.42-3.39 (4H, m), 3.28-3.26 (4H, m), 2.86 (3H, s), 1.42 (3H, t, $J=7.0$ Hz). HRMS (M-H) calcd for $\text{C}_{13}\text{H}_{19}\text{N}_4\text{O}_6\text{S}$: 359.10253; found: 359.1039.

10

Compound 83: N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-[4-(methylsulfonyl)-1-piperazinyl]-6-oxo-4-pyrimidinecarboxamide

15

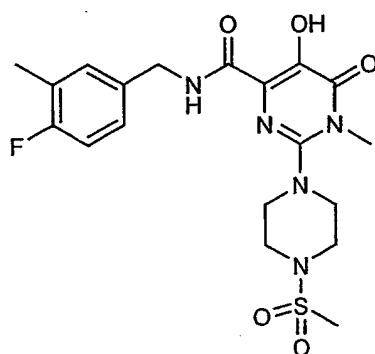


N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-[4-(methylsulfonyl)-1-piperazinyl]-6-oxo-4-pyrimidinecarboxamide was synthesized from 5-hydroxy-2-(4-methanesulfonyl-piperazin-1-yl)-1-methyl-6-oxo-1,6-dihydro-pyrimidine-4-

20

carboxylic acid ethyl ester and 4-fluorobenzylamine as described in example to yield the title compound as a white solid ^1H NMR (300 MHz, MeOD) δ : 8.88 (1H, t, $J=5.9$ Hz), 7.37 (2H, dd, $J=8.6, 5.3$ Hz), 7.05 (2H, t, $J=8.8$ Hz), 4.55 (2H, d, $J=6.2$ Hz), 3.55 (3H, s), 3.39-3.36 (4H, m), 3.26-3.23 (4H, m), 2.88 (3H, s). HRMS (M+H) calcd for $\text{C}_{18}\text{H}_{23}\text{FN}_5\text{O}_5\text{S}$: 440.14040; found: 440.1386.

Compound 84: N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-[4-(methylsulfonyl)-1-piperazinyl]-6-oxo-4-pyrimidinecarboxamide

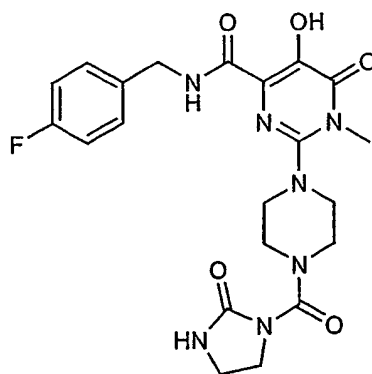


10

Pale lavender solid ^1H NMR (300 MHz, MeOD) δ : 8.83 (1H, bs), 7.21 (1H, d, $J=7.7$ Hz), 7.18-7.16 (1H, m), 6.96 (1H, t, $J=9.0$ Hz), 4.51 (2H, d, $J=4.0$ Hz), 3.55 (3H, s), 3.39-3.36 (4H, m), 3.26-3.23 (4H, m), 2.88 (3H, s), 2.24 (3H, d, $J=1.8$ Hz).

HRMS (M+H) calcd for $\text{C}_{19}\text{H}_{25}\text{FN}_5\text{O}_5\text{S}$: 454.15605; found: 454.1570.

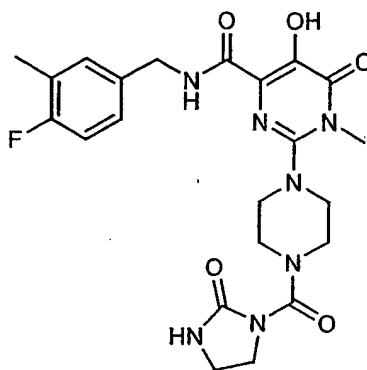
Compound 85: N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-[4-[(2-oxo-1-imidazolidinyl)carbonyl]-1-piperazinyl]-4-pyrimidinecarboxamide



20

White solid ^1H NMR (300 MHz, CDCl_3) δ : 11.74 (1H, s), 7.75 (1H, t, $J=5.7$ Hz), 7.32-7.27 (2H, m), 7.04 (2H, t, $J=8.6$ Hz), 4.82 (1H, s), 4.56 (2H, d, $J=6.2$ Hz), 3.87 (2H, t, $J=7.7$ Hz), 3.66-3.63 (4H, m), 3.55 (3H, s), 3.48 (2H, t, $J=7.7$ Hz), 3.12-3.09 (4H, m). HRMS (M+H) calcd for $\text{C}_{21}\text{H}_{25}\text{FN}_7\text{O}_5$: 474.19013; found: 474.1908.

Compound 86: N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-[4-[(2-oxo-1-imidazolidinyl)carbonyl]-1-piperazinyl]-4-pyrimidinecarboxamide

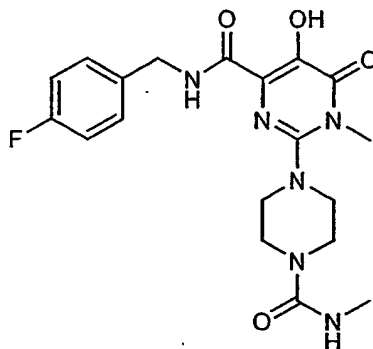


10

White solid ^1H NMR (300 MHz, CDCl_3) δ : 11.71 (1H, bs), 7.73 (1H, t, $J=6.0$ Hz), 7.14-7.02 (2H, m), 6.96 (1H, t, $J=8.8$ Hz), 5.50 (1H, bs), 4.51 (2H, d, $J=6.2$ Hz), 3.88 (2H, t, $J=7.9$ Hz), 3.66-3.64 (4H, m), 3.54 (3H, s), 3.51 (2H, t, $J=7.9$ Hz), 3.12-3.09 (4H, m). HRMS (M+H) calcd for $\text{C}_{22}\text{H}_{27}\text{FN}_7\text{O}_5\text{Na}$: 510.18772; found: 510.1884.

15

Compound 87: N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-[4-[(methylamino)carbonyl]-1-piperazinyl]-6-oxo-4-pyrimidinecarboxamide



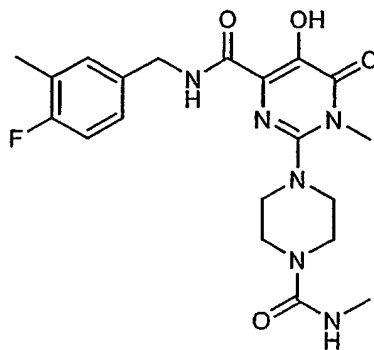
5

White solid ^1H NMR (300 MHz, MeOD) δ : 7.36 (2H, dd, $J=8.6, 5.3$ Hz), 7.05 (2H, t, $J=8.8$ Hz), 4.54 (2H, s), 3.56 (3H, s), 3.52 (4H, t, $J=4.9$ Hz), 3.11 (4H, t, $J=5.1$ Hz), 2.73 (3H, s). HRMS ($M+H$) calcd for $\text{C}_{19}\text{H}_{24}\text{FN}_6\text{O}_4$: 419.18432; found: 419.1858. CHN theoretical 1.0 mol H_2O : C, 52.29; H, 5.77; N, 19.26; F, 4.35;

10 found: C, 52.02; H, 5.72; N, 19.12; F, 4.13.

Compound 88: N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-[4-[(methylamino)carbonyl]-1-piperazinyl]-6-oxo-4-pyrimidinecarboxamide

15

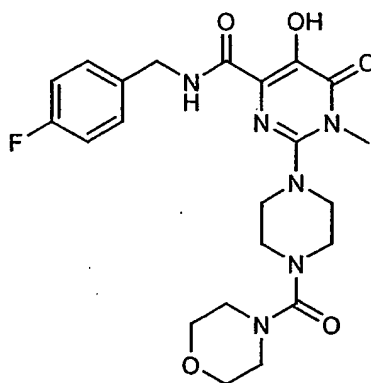


White solid ^1H NMR (300 MHz, MeOD) δ : 8.78 (1H, t, $J=5.8$ Hz), 7.20 (1H, d, $J=7.7$ Hz), 7.18-7.13 (1H, m), 6.96 (1H, t, $J=9.0$ Hz), 4.50 (2H, d, $J=4.0$ Hz), 3.55 (3H, s), 3.52 (4H, t, $J=4.9$ Hz), 3.11 (4H, t, $J=4.9$ Hz), 2.73 (3H, s), 2.23 (3H, d,

20

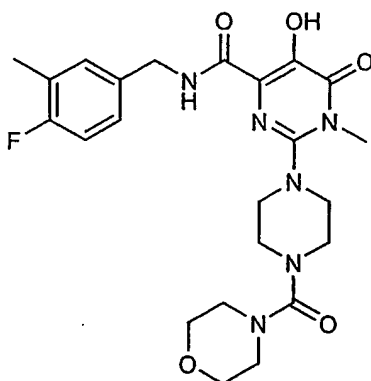
$J=1.8$ Hz). HRMS (M+H) calcd for $C_{20}H_{26}FN_6O_4$: 433.19997; found: 433.1990.
CHN theoretical 0.8 mol H_2O : C, 53.76; H, 6.00; N, 18.81; F, 4.25; found: C, 53.72; H, 5.61; N, 18.81; F, 4.19.

- 5 Compound 89: N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-[4-(4-morpholinylcarbonyl)-1-piperazinyl]-6-oxo-4-pyrimidinecarboxamide



- 10 White solid (0.064 g, 39% yield). 1H NMR (300 MHz, $CDCl_3$) δ : 7.72 (1H, t, $J=6.0$ Hz), 7.29 (2H, dd, $J=8.4, 5.5$ Hz), 7.04 (2H, t, $J=8.8$ Hz), 4.57 (2H, d, $J=6.2$ Hz), 3.71-3.67 (4H, m), 3.55 (3H, s), 3.45-3.42 (4H, m), 3.34-3.32 (4H, m), 3.06-3.03 (4H, m). HRMS (M-H) calcd for $C_{22}H_{26}N_6O_5F$: 473.19487; found: 473.1952.

- 15 Compound 90: N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-[4-(4-morpholinylcarbonyl)-1-piperazinyl]-6-oxo-4-pyrimidinecarboxamide



White solid (0.0648 g, 38% yield). ^1H NMR (300 MHz, CDCl_3) δ : 7.69 (1H, t, $J=5.8$ Hz), 7.14-7.08 (2H, m), 6.97 (1H, t, $J=8.8$ Hz), 4.52 (2H, d, $J=6.2$ Hz), 3.71-3.68 (4H, m), 3.56 (3H, s), 3.45-3.42 (4H, m), 3.34-3.31 (4H, m), 3.06-3.04 (4H, m),
5 2.27 (3H, s). HRMS (M-H) calcd for $\text{C}_{23}\text{H}_{28}\text{N}_6\text{O}_5\text{F}$: 487.21052; found: 487.2094.
CHN theoretical 0.4 mol H_2O , 1.0 mol TFA: C, 49.25; H, 5.09; N, 13.78; F, 12.46;
found: C, 49.08; H, 5.08; N, 13.61; F, 12.23.

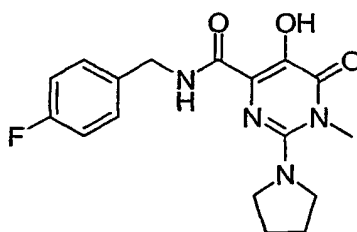
EXAMPLE 12

10 Preparation of compounds 91-101

Compounds 91-101 were prepared according to the methods described above.

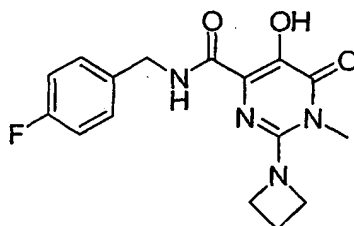
Compound 91: N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-
2-(1-pyrrolidinyl)-4-pyrimidinecarboxamide

15



HPLC 100% AP (RT, 2.44 min); MS (ESI) m/z 347 (M+H), 345 (M-H);
HRMS (ESI) calcd for $\text{C}_{17}\text{H}_{20}\text{N}_4\text{O}_3\text{F}$ (M+H) 347.1519, found 347.1520 (δ +0.2
20 ppm); ^1H NMR (CDCl_3 , 300 MHz) δ ppm 1.91 (4H, m), 3.45 (4H, t, $J=6.6$ Hz), 3.49
(3H, s), 4.56 (2H, d, $J=6$ Hz), 7.04 (2H, m), 7.30 (2H, m), 7.80 (1H, br), 11.34 (1H, s)

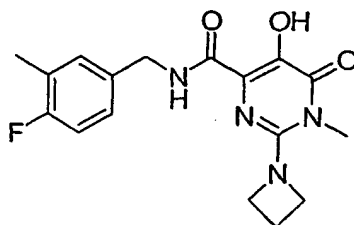
Compound 92: 2-(1-azetidinyl)-N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-4-pyrimidinecarboxamide



5

HPLC 98%AP (RT, 2.23 min); HRMS (ESI) calcd for $C_{16}H_{18}N_4O_3F$ (M+H) 333.1363, found 333.1360 (δ -0.6 ppm); 1H NMR ($CDCl_3$, 500MHz) δ ppm 2.30 (2H, qi, $J=7.5$ Hz), 3.37 (3H, s), 4.05 (4H, t, $J=7.5$ Hz), 4.55, (2H, q, $J=6.5$ Hz), 7.02 (2H, td, $J=7, 2$ Hz), 7.29 (2H, m), 7.82 (1H, br), 11.35 (1H, s); ^{13}C NMR ($CDCl_3$, 125.8MHz) δ ppm 16.2, 31.1, 42.3, 52.6, 115.7, 115.8, 125.3, 125.8, 129.3, 129.4, 133.4, 133.4, 142.4, 149.7, 159.5, 161.4, 163.4, 168.6; UV (MeOH) λ_{max} 245nm ($\epsilon 1.03 \times 10^4$), 337 ($\epsilon 5.0 \times 10^3$); Elemental Analysis calcd for $C_{16}H_{17}N_4O_3F \cdot 1/5H_2O$; C57.20, H5.22, N16.68; found C57.43, H5.77, N16.86.

15 Compound 93: 2-(1-azetidinyl)-N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-4-pyrimidinecarboxamide



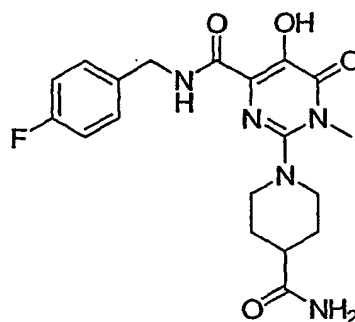
20

HPLC 96%AP (RT, 2.44 min); HRMS (ESI) calcd for $C_{17}H_{20}N_4O_3F$ (M+H) 347.1519, found 347.1515 (δ -1.3 ppm); 1H NMR ($CDCl_3$, 500MHz) δ ppm 2.25 (3H, s), 2.30 (2H, qi, $J=7.5$ Hz), 3.37 (3H, s), 4.05 (4H, t, $J=7.5$ Hz), 4.50, (2H, q, $J=6.5$ Hz), 6.95 (1H, t, $J=9$ Hz), 7.07-7.13 (2H, m), 7.81 (1H, br), ~11 (1H, br); ^{13}C NMR ($CDCl_3$, 125.8MHz) δ ppm 14.5, 14.5, 16.2, 31.1, 42.4, 52.7, 115.2, 115.4,

125.3, 125.4, 125.5, 126.5, 126.5, 130.8, 130.9, 133.0, 133.1, 142.3, 149.7, 159.6, 159.9, 161.9, 168.5; UV (MeOH) λ_{max} 245nm (ϵ 8.95 $\times 10^3$), 336 (ϵ 4.43 $\times 10^3$); Elemental Analysis calcd for C₁₇H₁₉N₄O₃F.1/2H₂O, C57.46, H5.67, N15.77; found C57.87, H5.62, N15.46.

5

Compound 94: 2-[4-(aminocarbonyl)-1-piperidiny]-N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-4-pyrimidinecarboxamide

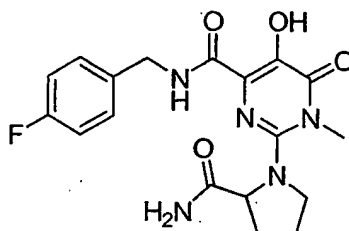


10

HPLC 96%AP (RT, 2.44 min); HRMS (ESI) calcd for for C₁₇H₂₀N₄O₃F (M+H) 347.1519, found 347.1515 (δ -1.3 ppm); ¹H NMR (CDCl₃, 500MHz) δ ppm 2.25 (3H, s), 2.30 (2H, qi, J=7.5Hz), 3.37 (3H, s), 4.05 (4H, t, J=7.5Hz), 4.50, (2H, q, J = 6.5Hz), 6.95 (1H, t, J=9Hz), 7.07-7.13 (2H, m), 7.81 (1H, br), ~11 (1H, br); ¹³C NMR (CDCl₃, 125.8MHz) δ ppm 14.5, 14.5, 16.2, 31.1, 42.4, 52.7, 115.2, 115.4, 125.3, 125.4, 125.5, 126.5, 126.5, 130.8, 130.9, 133.0, 133.1, 142.3, 149.7, 159.6, 159.9, 161.9, 168.5; UV (MeOH) λ_{max} 245nm (ϵ 8.95 $\times 10^3$), 336 (ϵ 4.43 $\times 10^3$); Elemental Analysis calcd for C₁₇H₁₉N₄O₃F.1/2H₂O, C57.46, H5.67, N15.77; found C57.87, H5.62, N15.46.

20

Compound 95: 2-[(2S)-2-(aminocarbonyl)-1-pyrrolidinyl]-N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-4-pyrimidinecarboxamide



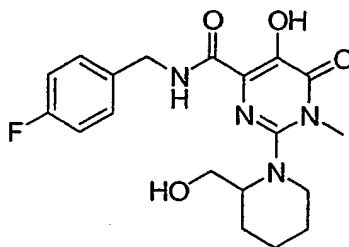
5

Yield 67% (white solid, trituration with Et₂O); HPLC ~95%AP (RT, 1.80 min); HRMS (ESI) calcd for C₁₈H₂₁N₅O₄F (M+H) 390.1578, found 390.1570 (δ - 1.9 ppm); ¹H NMR (CDCl₃, 300 MHz) δ ppm 1.97 (2H, m), 2.21 (1H, m), 2.40 (1H, m), 3.26 (1H, br.t, J=9Hz), 3.56 (3H, s), 3.76 (1H, m), 4.46-4.60 (3H, m), 5.19 (1H, br.s), 5.91 (1H, br.s), 7.02 (2H, t, J=8.4Hz), 7.30 (2H, m), 7.86 (1H, br), 11.47 (1H, s).

10

Compound 96: 4-pyrimidineN-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-2-[2-(hydroxymethyl)-1-piperidinyl]-1-methyl-6-oxo-carboxamide

15

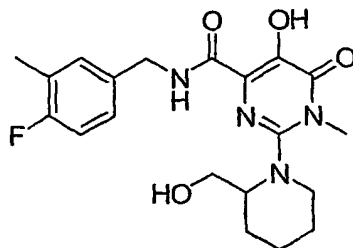


¹H NMR (300 MHz, MeOD) δ : 8.90 (1H, bs), 7.39-7.35 (2H, m), 7.05 (2H, t, J=8.8 Hz), 4.72 (1H, dd, J=12.6, 2.7 Hz), 4.56 (2H, s), 4.57-4.49 (1H, m), 3.65-3.57 (1H, m), 3.49 (3H, s), 3.49-3.43 (1H, m), 3.11-3.02 (1H, m), 2.04-1.92 (3H, m), 1.72-1.58 (3H, m). HRMS (M+H) calcd for C₁₉H₂₄FN₄O₄: 391.17817; found: 391.1783. CHN theoretical for 1.3 mol TFA and 0.55 mol H₂O: C, 47.29; H, 4.67; N, 10.21; F, 16.97; found: C, 47.69; H, 4.36; N, 10.23; F, 16.86.

20

25

Compound 97: N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-2-[2-(hydroxymethyl)-1-piperidinyl]-1-methyl-6-oxo-4-pyrimidinecarboxamide



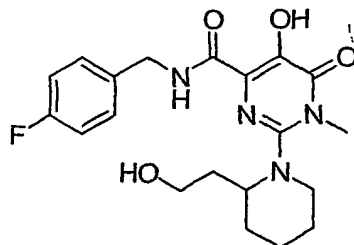
5

^1H NMR (300 MHz, MeOD) δ : 8.90 (1H, bs), 7.22-7.14 (2H, m), 6.96 (1H, t, $J=9.0$ Hz), 4.75 (1H, dd, $J=12.8, 2.9$ Hz), 4.51 (2H, s), 3.65-3.57 (1H, m), 3.48 (3H, s), 3.48-3.44 (1H, m), 3.11-3.03 (1H, m), 2.23 (3H, s), 2.04-1.91 (3H, m), 1.73-1.57 (3H, m). HRMS (M+H) calcd for $\text{C}_{20}\text{H}_{26}\text{FN}_4\text{O}_4$: 405.19382; found: 405.1958.

10 CHN theoretical for 1.4 mol TFA and 0.3 mol H_2O : C, 48.09; H, 4.78; N, 9.84; F, 17.35; found: C, 48.41; H, 4.47; N, 9.63; F, 17.18.

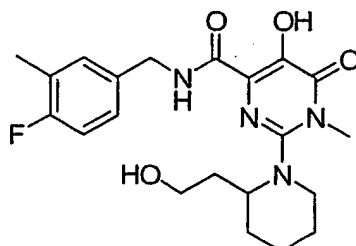
Compound 98: N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-2-[2-(2-hydroxyethyl)-1-piperidinyl]-1-methyl-6-oxo-4-pyrimidinecarboxamide

15



^1H NMR (300 MHz, CDCl_3) δ : 9.18-9.10 (1H, m), 8.65-8.60 (1H, m), 8.27 (1H, t, $J=6.4$ Hz), 7.31 (2H, dd, $J=8.6, 5.3$ Hz), 6.99 (2H, t, $J=8.6$ Hz), 4.75-4.67 (1H, m), 4.52-4.49 (2H, m), 4.41-4.34 (1H, m), 3.32 (3H, s), 3.36-3.22 (2H, m), 2.95-2.83 (1H, m), 2.35-2.25 (1H, m), 2.16-2.07 (1H, m), 1.94-1.90 (2H, d, $J=11.71$ Hz), 1.80-1.66 (3H, m), 1.56-1.46 (1H, m). HRMS (M-H) calcd for $\text{C}_{20}\text{H}_{24}\text{FN}_4\text{O}_4$: 403.17816; found: 403.1780. CHN theoretical for 1.4 mol TFA and 0.4 mol H_2O : C, 47.94; H, 4.80; N, 9.81; F, 17.29; found: C, 47.87; H, 4.62; N, 9.61; F, 17.58.

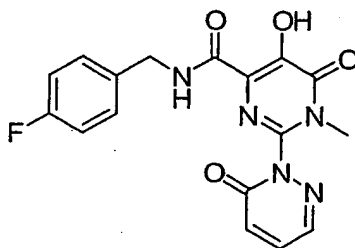
Compound 99: N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-2-[2-(2-hydroxyethyl)-1-piperidinyl]-1-methyl-6-oxo-4-pyrimidinecarboxamide



5

^1H NMR (300 MHz, CDCl_3) δ : 9.14-9.09 (1H, m), 8.57-8.53 (1H, m), 8.24 (1H, t, $J=6.4$ Hz), 7.16-7.11 (2H, m), 6.92 (1H, t, $J=8.8$ Hz), 4.78-4.70 (1H, m), 4.47 (2H, t, $J=5.1$ Hz), 4.42-4.36 (1H, m), 3.34 (3H, s), 3.34-3.19 (2H, m), 2.92-2.84 (1H, m), 2.35-2.25 (1H, m), 2.23 (3H, s), 2.18-2.09 (1H, m), 1.92 (2H, d, $J=12.1$ Hz), 1.82-1.67 (3H, m), 1.57-1.46 (1H, m). HRMS (M-H) calcd for $\text{C}_{21}\text{H}_{26}\text{FN}_4\text{O}_4$: 417.19381; found: 417.1930. CHN theoretical for 1.5 mol TFA and 0.5 mol H_2O : C, 48.16; H, 4.97; N, 9.36; F, 17.46; found: C, 48.43; H, 4.81; N, 9.34; F, 17.46.

15 Compound 100: N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(6-oxo-1(6H)-pyridazinyl)-4-pyrimidinecarboxamide



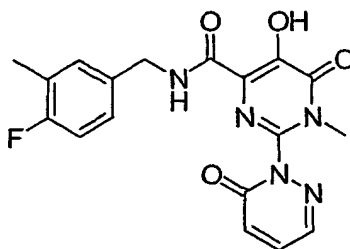
20

HPLC 98% AP (RT, 1.98 min); HRMS (ESI) calcd for $\text{C}_{17}\text{H}_{13}\text{N}_5\text{O}_4\text{F}$ (M-H) 370.0952, found 370.0937 (δ -3.9 ppm); ^1H NMR (CDCl_3 , 500MHz) δ ppm 3.37 (3H, s), 4.52, (2H, q, $J=6$ Hz), 6.99-7.03 (2H, m), 7.06 (1H, dd, $J=10, 1.5$ Hz), 7.25-7.28 (2H, m), 7.36 (1H, dd, $J=10, 4$ Hz), 7.66 (1H, br.t, $J=6$ Hz), 7.93 (1H, dd, $J=4, 1.5$ Hz), 12.49 (1H, s); ^{13}C NMR (CDCl_3 , 125.8MHz) δ ppm 31.1, 42.6, 115.71,

115.87, 124.6, 129.81, 129.87, 131.6, 132.7, 133.1, 138.1, 140.1, 149.2, 158.2, 159.1, 161.47, 163.43, 167.6; UV (MeOH) λ_{max} 251nm (ϵ 1.01x10⁴), 297 (ϵ 1.13x10⁴); Elemental Analysis calcd for C₁₇H₁₄N₅O₄F, C54.98, H3.80, N18.86; found C54.72, H3.64, N18.86.

5

Compound 101: N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(6-oxo-1(6H)-pyridazinyl)-4-pyrimidinecarboxamide

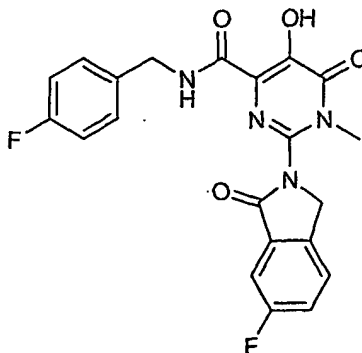


10

HPLC 96% AP (RT, 2.16 min); HRMS (ESI) calcd for for C₁₈H₁₅N₅O₄F (M-H) 384.1108, found 384.1112 (δ +1.0 ppm); ¹H NMR (CDCl₃, 500MHz) δ ppm 2.23 (3H, s), 3.37 (3H, s), 4.48, (2H, q, J = 6Hz), 6.94 (1H, t, J=9Hz), 7.06 (1H, dd, J=9.5, 1.5Hz), 7.07 (1H, m), 7.11 (1H, m), 7.37 (1H, dd, J= 9.5, 3.5Hz), 7.63 (1H, br.t, J=6Hz), 7.94 (1H, dd, J=3.5, 1.5Hz), 12.53 (1H, s); ¹³C NMR (CDCl₃, 125.8MHz) δ ppm 14.58, 14.61, 31.1, 42.7, 115.2, 115.4, 124.6, 125.4, 125.5, 127.0, 127.1, 131.31, 131.35, 131.5, 132.3, 132.4, 133.1, 138.1, 140.1, 149.2, 158.2, 159.1, 160.02, 161.98, 167.6; UV (MeOH) λ_{max} 251nm (ϵ 8.75x10³), 297 (ϵ 9.80x10³); Elemental Analysis calcd for C₁₈H₁₆N₅O₄F, C56.10, H4.18, N18.17; found C55.17, H3.72, N17.91

20

Compound 102: 2-(6-fluoro-1,3-dihydro-1-oxo-2H-isoindol-2-yl)-N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-4-pyrimidinecarboxamide



5

HPLC 100% AP (RT, 2.51 min); HRMS (ESI) calcd for $C_{21}H_{17}N_4O_4F_2$ (M+H) 427.1218, found 427.1212 (δ -0.6 ppm); 1H NMR ($CDCl_3$, 300 MHz) δ ppm 3.59 (3H, s), 4.59 (2H, q, J = 6.2 Hz), 4.91 (2H, s), 7.02 (2H, t, J = 8.7 Hz), 7.27-7.59 (5H, m), 12.17 (1H, s)

10

BIOLOGICAL ACTIVITY

HIV-Integrase Inhibition Activity

- 15 To evaluate *in-vitro* activity against HIV-integrase, 5 pmole of biotin labeled substrate DNA was bound to 100 μ g of Streptavidin coated PVT SPA beads (Amersham Pharmacia Biotech). Recombinant integrase (0.26 ng) was incubated with the beads for 90 min at 37 °C. Unbound enzyme was removed by washing the complex followed by addition of inhibitors and 0.1 fmol of P33 labeled target DNA.
- 20 The reaction was stopped by adding EDTA to a final concentration of 10 mM. Samples were counted in TopCountNXT (Packard) and the CPM was used as a measure of integration. The reaction condition was as described in A. Engelman and R. Craigie, J. Virol. 69, 5908-5911 (1995). The sequences of substrate and target DNA were described in Nucleic Acid Research 22, 1121-1122 (1994). Results are
- 25 shown in the table. Activity equal to A refers to a compound having $IC_{50} = 0.003$ to

0.10 μM while B and C denote compounds the following $\text{IC}_{50} = 0.1$ to $1.0 \mu\text{M}$ and $\text{IC}_{50} \geq 1.0 \mu\text{M}$ respectively.

Compound	Activity
1	A
2	A
3	B
4	A
5	A
6	A
7	C
8	A
9	A
10	A
11	B
12	A
13	A
14	A
15	A
16	C
17	A
18	A
19	A
20	A
21	C
22	A
23	A
24	A
25	C
26	C
27	C

Compound	Activity
28	B
29	A
30	B
31	C
32	B
33	B
34	B
35	B
36	B
37	B.
38	C
39	C
40	C
41	C
42	B
43	B
44	C
45	B
46	B
47	B
48	B
49	C
50	C
52	A
53	A
54	A
55	A
56	A
57	A
58	B

Compound	Activity
59	A
60	A
61	B
62	A
63	A
64	A
65	A
66	A
67	A
68	A
69	B
70	A
71	A
72	A
73	A
74	A
75	A
76	A
77	A
78	A
79	A
80	A
84	
85	A
86	A
87	A
88	A
91	A
92	A
93	A

Compound	Activity
94	A
95	A
96	B
97	A
98	A
99	A
100	B
101	A
102	A

Inhibition of HIV replication

A recombinant NL-Rluc virus was constructed in which a section of the nef
5 gene from NL4-3 was replaced with the *Renilla* Luciferase gene. The NL-RLuc virus
was prepared by co-transfection of two plasmids, pNLRLuc and pVSVenv. The
pNLRLuc contains the NL-Rluc DNA cloned into pUC18 at the *PvuII* site, while the
pVSVenv contains the gene for VSV G protein linked to an LTR promoter.
Transfections were performed at a 1:3 ratio of pNLRLuc to pVSVenv on 293T cells
10 using the LipofectAMINE PLUS kit from Invitrogen (Carlsbad, CA) according to
manufactures instruction, and the pseudotype virus generated was titered in MT-2
cells.

Susceptibility of viruses to compounds was determined by incubation in the
presence of serial dilutions of the compound. The 50% effective concentration
15 (EC₅₀) was calculated by using the exponential form of the median effect equation
where $(Fa) = 1/[1 + (ED_{50}/\text{drug conc.})^m]$ (Johnson VA, Byington RT. Infectivity
Assay. In *Techniques in HIV Research*. ed. Aldovini A, Walker BD. 71-76. New
York: Stockton Press.1990). The anti-viral activity of compounds was evaluated
under three serum conditions, 10% FBS, 15mg/ml human serum albumin/10% FBS
20 or 40% human serum/5% FBS, and the results from at least 2 experiments were used
to calculate the EC₅₀ values. Results are shown in the table. Activity equal to A

refers to a compound having $EC_{50} = 0.003$ to $0.10 \mu\text{M}$ while B and C denote compounds the following $EC_{50} = 0.1$ to $1.0 \mu\text{M}$ and $EC_{50} \geq 1.0 \mu\text{M}$ respectively.

Compound	Activity
1	B
2	A
3	C
4	B
5	B
6	B
8	B
9	A
10	B
11	B
12	A
13	A
14	B
15	B
17	C
18	A
19	A
20	B
22	B
23	B
24	B
28	C
29	B
32	C
35	C
36	C
37	C
42	C

Compound	Activity
43	C
45	C
46	C
47	C
48	C
52	B
53	A
54	A
55	A
56	A
57	A
58	C
59	C
60	C
61	B
62	B
63	B
65	A
66	A
67	B
68	B
69	C
70	B
71	B
72	A
73	B
74	A
75	A
76	A
77	A

Compound	Activity
78	A
79	A
80	B
85	B
86	A
87	B
88	A
91	A
92	A
94	B

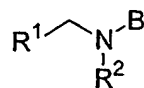
5

Although the invention has been described with respect to specific aspects, those skilled in the art will recognize that other aspects not specifically described herein are intended to be within the scope of the claims which follow.

CLAIMS

1. A compound having the formula

5



(I)

wherein:

10 R^1 is $-\text{C}_1\text{-C}_6$ alkyl, $\text{C}_3\text{-C}_6$ cycloalkyl, -aryl, $-\text{C}_1\text{-C}_6$ alkyl-aryl, or $-\text{C}_1\text{-C}_6$ alkyl-diaryl;

R^2 is -H, -OH, $-\text{C}_1\text{-C}_6$ alkyl unsubstituted or substituted with 1-3 R^3 , or $-\text{OR}^4$;

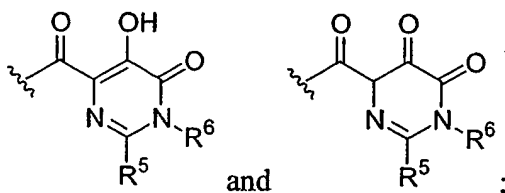
Each R^3 is independently selected from the group consisting of -H, -halo, -CN, $-\text{C}_1\text{-C}_2$ perfluoroalkyl, $-\text{C}_1\text{-C}_6$ alkyl, $-\text{C}_1\text{-C}_2$ perfluoroalkoxy, and $-\text{C}_1\text{-C}_6$ alkoxy;

15

R^4 is $-\text{C}_1\text{-C}_6$ alkyl or $-\text{C}_3\text{-C}_6$ cycloalkyl;

B is selected from the group consisting of

20



R^5 is $-\text{NR}^7\text{R}^8$; or Het wherein the bond attaching R^5 is made to a heteroatom in said Het;

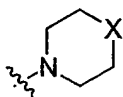
25

R^6 is -H, $-\text{C}_1\text{-C}_6$ alkyl unsubstituted or substituted with 1-3 R^3 ; and

R^7 and R^8 are independently -H or $-\text{C}_1\text{-C}_6$ alkyl;

or pharmaceutically acceptable enantiomer, diastereomer, salt, solvate or prodrug thereof.

- 5 2. The compound of claim 1 wherein the aryl is substituted with 1-3 R^3 .
3. The compound of claim 1 wherein R^1 is -aryl, - C_1 - C_6 alkyl-aryl, or - C_1 - C_6 alkyl diaryl.
- 10 4. The compound of claim 3 wherein R^1 is -aryl, - C_1 - C_6 alkyl-phenyl, or - C_1 - C_6 alkyl-diphenyl.
5. The compound of claim 4 wherein the aryl is selected from the group consisting of phenyl, naphthyl, furanyl, thienyl, indolyl, and benzimidazolyl, and
- 15 aryl and phenyl are unsubstituted or substituted with 1-3 R^3 .
6. A compound of claim 4 wherein R^1 is aryl selected from the group consisting of naphthyl, 2-methyl-5-furanyl, 2-thienyl, 3-indolyl, and 2-benzimidazolyl.
- 20 7. A compound of claim 4 wherein R^1 is phenyl unsubstituted or substituted with 1-3 R^3 .
8. A compound of claim 7 wherein R^1 is 3-fluorophenyl, 4-fluorophenyl,
- 25 2,4-difluorophenyl, 3,4-difluorophenyl, 3-methyl-4-fluorophenyl, 3-chloro-4-fluorophenyl, 3-chlorophenyl, 4-chlorophenyl, 3,4-dichlorophenyl, 3,5-dichlorophenyl, 3-methylphenyl, 4-methylphenyl, or 3,4-dimethylphenyl.
9. A compound of claim 1 where R^2 is -H, methyl, -OH, or methoxy.
- 30 10. A compound of claim 9 where R^2 is -H.

11. A compound of claim 1 wherein R⁶ is -C₁-C₆ alkyl.
12. A compound of claim 11 wherein R⁶ is methyl.
- 5 13. A compound of claim 1 wherein R⁵ is heteroaryl.
14. A compound of claim 13 wherein R⁵ is selected from the group consisting of fluorooxindole and methyltriazole.
- 10 15. A compound of claim 1 wherein R⁵ is a 4 to 7 membered monocyclic heterocycle.
16. A compound of claim 15 wherein R⁵ is selected from the group consisting of [1,2]thiazinane 1,1-dioxide and , wherein X is a bond, O, S, NH, N-methyl, N-propyl, N-acetyl, N-(2-hydroxyethyl), N-pyridinyl, N-pyrazinyl, N-SO₂Me, CH₂, and CHOH.
- 15
17. A compound of claim 1 selected from the group consisting of
 - 20 N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
 - N-[(3,4-dichlorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
 - 25 N-[(3-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
 - N-[(4-chlorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
 - 30 N-[(3-chlorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
 - 35 N-[3-(4-fluorophenyl)propyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;

- N-[3-(3,4-dichlorophenyl)propyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 5 1,6-dihydro-5-hydroxy-1-methyl-N-[(4-methylphenyl)methyl]-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 1,6-dihydro-5-hydroxy-1-methyl-N-[(3-methylphenyl)methyl]-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 10 1,6-dihydro-5-hydroxy-N-[(4-methoxyphenyl)methyl]-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 1,6-dihydro-5-hydroxy-N-[(3-methoxyphenyl)methyl]-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 15 N-[(3,5-dichlorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 20 N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 25 N-[(3-fluoro-4-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- N-[(3,4-difluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 30 N-[2-(4-fluorophenyl)ethyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- N-[(2,4-difluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 35 N-[(3-chloro-4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 40 N-[(3,4-dimethylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 1,6-dihydro-5-hydroxy-1-methyl-6-oxo-N-(1-phenylethyl)-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 45 N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(4-morpholinyl)-6-oxo-4-pyrimidinecarboxamide;

- N-[[4-fluoro-3-(trifluoromethyl)phenyl]methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 5 N-[2-(3-fluorophenyl)ethyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- N-[(2,4-dimethoxyphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 10 1,6-dihydro-5-hydroxy-1-methyl-N-[(5-methyl-2-furanyl)methyl]-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 1,6-dihydro-5-hydroxy-N-[(2-methoxyphenyl)methyl]-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 15 N-(2,2-diphenylethyl)-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 20 N-[2-(4-chlorophenyl)ethyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 1,6-dihydro-5-hydroxy-N-[2-(4-methoxyphenyl)ethyl]-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 25 N-[(2-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-N-[[3-(trifluoromethyl)phenyl]methyl]-4-pyrimidinecarboxamide;
- 30 N-[2-(3-chlorophenyl)ethyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 35 1,6-dihydro-5-hydroxy-1-methyl-6-oxo-N-(2-phenylethyl)-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 1,6-dihydro-5-hydroxy-1-methyl-6-oxo-N-[(1S)-1-phenylethyl]-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 40 1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-N-[[4-(trifluoromethyl)phenyl]methyl]-4-pyrimidinecarboxamide;
- 45 N-[1-(4-fluorophenyl)ethyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;

- N-(2-furanylmethyl)-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 5 1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-N-(2-thienylmethyl)-4-pyrimidinecarboxamide;
- N-(1,2-diphenylethyl)-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 10 1,6-dihydro-5-hydroxy-1-methyl-6-oxo-N-(phenylmethyl)-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 15 N-[(2-chlorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 1,6-dihydro-5-hydroxy-1-methyl-N-[(2-methylphenyl)methyl]-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 20 1,6-dihydro-5-hydroxy-N-[2-(2-methoxyphenyl)ethyl]-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 1,6-dihydro-5-hydroxy-N-[2-(3-methoxyphenyl)ethyl]-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 25 1,6-dihydro-5-hydroxy-1-methyl-N-[2-(4-methylphenyl)ethyl]-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-N-[(3,4,5-trimethoxyphenyl)methyl]-4-pyrimidinecarboxamide;
- 30 N-[(2,5-dichlorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 35 1,6-dihydro-5-hydroxy-N-[(1R,2S)-2-hydroxy-1,2-diphenylethyl]-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 40 N-[(2,4-dichlorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- N-[[3,5-bis(trifluoromethyl)phenyl]methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 45 N-[(2,5-difluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;

- 1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-N-[[2-(trifluoromethyl)phenyl]methyl]-4-pyrimidinecarboxamide;
- 5
- 1,6-dihydro-5-hydroxy-1-methyl-6-oxo-N-(1-phenylpropyl)-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 10
- 4-pyrimidinecarboxamide, N-[1-(4-chlorophenyl)ethyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 15
- N-[(3,5-difluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-N-[[3-(trifluoromethoxy)phenyl]methyl]-4-pyrimidinecarboxamide;
- 20
- 1,6-dihydro-5-hydroxy-1-methyl-N-[1-(1-naphthalenyl)ethyl]-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 25
- N-[1-(4-bromophenyl)ethyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 1,6-dihydro-5-hydroxy-1-methyl-N-[1-(2-naphthalenyl)ethyl]-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 30
- 1,6-dihydro-5-hydroxy-1-methyl-N-[1-(4-methylphenyl)ethyl]-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 1,6-dihydro-5-hydroxy-N-[1-(3-methoxyphenyl)ethyl]-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 35
- 1,6-dihydro-5-hydroxy-N-[1-(4-methoxyphenyl)ethyl]-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 40
- 1,6-dihydro-5-hydroxy-N-[2-(1H-indol-3-yl)ethyl]-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 1,6-dihydro-5-hydroxy-1-methyl-6-oxo-N-[(1S)-1-phenylpropyl]-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;
- 45
- N-(1H-benzimidazol-2-ylmethyl)-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)-4-pyrimidinecarboxamide;

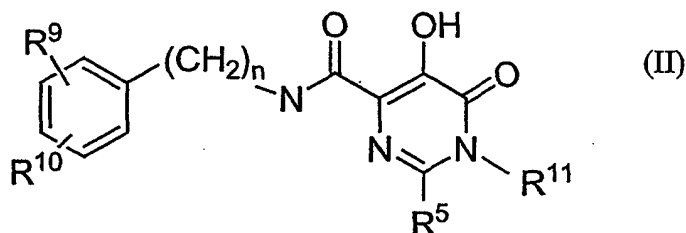
- N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(3-methyl-1H-1,2,4-triazol-1-yl)-6-oxo-4-pyrimidinecarboxamide;
- 5 N-[(3,4-dichlorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(3-methyl-1H-1,2,4-triazol-1-yl)-6-oxo-4-pyrimidinecarboxamide;
- N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(3-methyl-1H-1,2,4-triazol-1-yl)-6-oxo-4-pyrimidinecarboxamide;
- 10 N-[(3,4-dichlorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(4-morpholinyl)-6-oxo-4-pyrimidinecarboxamide;
- N-[(3-fluoro-4-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(4-morpholinyl)-6-oxo-4-pyrimidinecarboxamide;
- 15 N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(4-morpholinyl)-6-oxo-4-pyrimidinecarboxamide;
- N-[(3-chloro-4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(4-morpholinyl)-6-oxo-4-pyrimidinecarboxamide;
- 20 N-[(3,4-dimethylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(4-morpholinyl)-6-oxo-4-pyrimidinecarboxamide;
- 25 N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(5-methyl-1H-1,2,4-triazol-1-yl)-6-oxo-4-pyrimidinecarboxamide;
- N-[(3,4-dichlorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-(5-methyl-1H-1,2,4-triazol-1-yl)-6-oxo-4-pyrimidinecarboxamide;
- 30 N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-2-[4-(2-hydroxyethyl)-1-piperazinyl]-1-methyl-6-oxo-4-pyrimidinecarboxamide;
- N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-2-[4-(2-hydroxyethyl)-1-piperazinyl]-1-methyl-6-oxo-4-pyrimidinecarboxamide;
- 35 2-(4-acetyl-1-piperazinyl)-N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-4-pyrimidinecarboxamide;
- 40 2-(4-acetyl-1-piperazinyl)-N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-4-pyrimidinecarboxamide;
- N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(4-propyl-1-piperazinyl)-4-pyrimidinecarboxamide;
- 45 N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(4-propyl-1-piperazinyl)-4-pyrimidinecarboxamide;

- N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(1-piperazinyl)-4-pyrimidinecarboxamide;
- 5 N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(1-piperazinyl)-4-pyrimidinecarboxamide;
- N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-2-(4-hydroxy-1-piperidinyl)-1-methyl-6-oxo-4-pyrimidinecarboxamide;
- 10 N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-2-(4-hydroxy-1-piperidinyl)-1-methyl-6-oxo-4-pyrimidinecarboxamide;
- 2-(6-fluoro-1,3-dihydro-1-oxo-2H-isoindol-2-yl)-N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-4-pyrimidinecarboxamide;
- 15 N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-[4-(2-pyridinyl)-1-piperazinyl]-4-pyrimidinecarboxamide;
- 20 N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-[4-(2-pyridinyl)-1-piperazinyl]-4-pyrimidinecarboxamide;
- N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-[4-(2-pyrimidinyl)-1-piperazinyl]-4-pyrimidinecarboxamide;
- 25 N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-[4-(2-pyrimidinyl)-1-piperazinyl]-4-pyrimidinecarboxamide;
- N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-[4-(methylsulfonyl)-1-piperazinyl]-6-oxo-4-pyrimidinecarboxamide;
- 30 N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-[4-(methylsulfonyl)-1-piperazinyl]-6-oxo-4-pyrimidinecarboxamide;
- N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-(1-pyrrolidinyl)-4-pyrimidinecarboxamide;
- 35 N-[(4-fluoro-3-methylphenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-[2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)ethyl]-4-pyrimidinecarboxamide;
- 40 N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-[3-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)propyl]-4-pyrimidinecarboxamide;
- 45

N-[(4-fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-[2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)ethyl]-4-pyrimidinecarboxamide;
and

- 5 N-[(3,4-dichlorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-6-oxo-2-[2-(tetrahydro-1,1-dioxido-2H-1,2-thiazin-2-yl)ethyl]-4-pyrimidinecarboxamide.

18. A compound having the formula:



10

wherein,

n is 1 to 4;

R⁵ is Het wherein the bond attaching R⁵ is made to a heteroatom in said Het;

15

R⁹ and R¹⁰ are each independently H, halo, C₁₋₆ alkyl, C₁₋₆ alkoxy or CF₃; and

R¹¹ is C₁₋₆ alkyl;

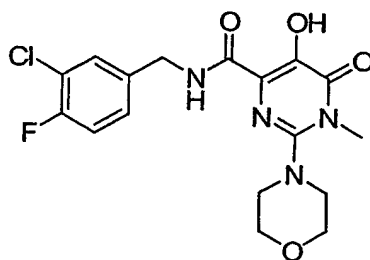
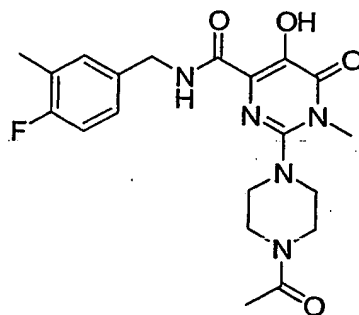
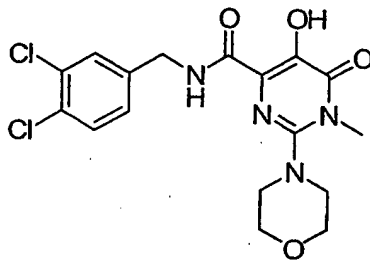
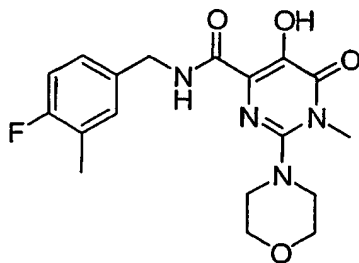
- 20 or a pharmaceutically acceptable enantiomer, diastereomer, salt, solvate or prodrug thereof.

19. The compound of claim 18 wherein R⁵ is a C₄₋₇ monocyclic heterocycle.

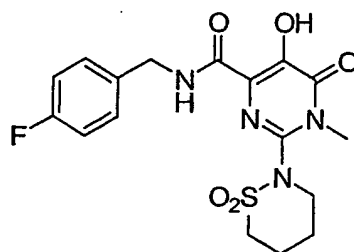
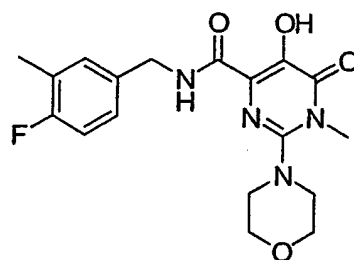
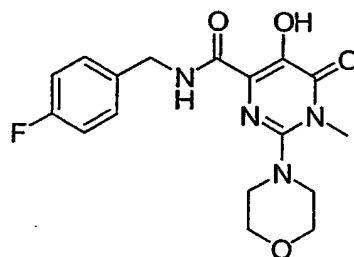
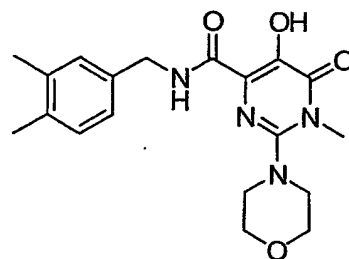
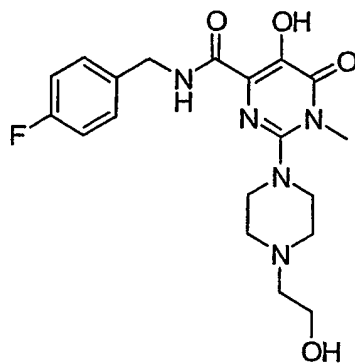
- 25 20. The compound of claim 19 wherein the heterocycle is substituted with at least one of C₁₋₆ alkyl, C₁₋₆ alkoxy, C₁₋₆ hydroxyalkyl, C₁₋₆ alkyl carbonyl, C₆₋₁₀ aryl, C₇₋₁₄ alkyl, aryl, C₁₋₆ alkylsulfonyl, C₆₋₁₀ arylsulfonyl, C₁₋₆ alkylurea, C₃₋₇ cycloalkylurea, carbonyl, carboxamide or hydroxyl.

- 30 21. The compound of claim 18 having one of the following structures;

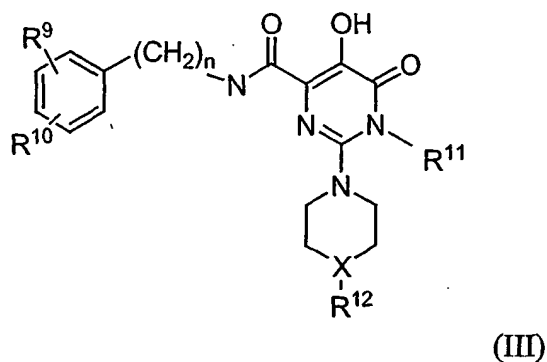
130



131



22. A compound having the formula:



5 wherein n is 1 to 4;

R^9 and R^{10} are each independently H, halo, C_{1-6} alkyl, C_{1-6} alkoxy or CF_3 ;

X is C, O or N;

R^{11} is C_{1-6} alkyl; and

10 R^{12} is H, C_{1-6} alkyl, C_{1-6} alkoxy, C_{1-6} hydroxyalkyl, C_{1-6} alkyl carbonyl, C_{6-10} aryl, C_{7-14} alkyl; aryl, C_{1-6} alkylsulfonyl, C_{6-10} arylsulfonyl, C_{1-6} alkylurea, C_{3-7} cycloalkylurea, carbonyl, carboxamide or hydroxyl;

said R^{12} being present only when X is C or N;

or a pharmaceutically acceptable enantiomer, diastereomer, salt, solvate or prodrug thereof.

15

23. The compound of claim 22 wherein m is 1.

24. The compound of claim 22 wherein R_9 is halo.

20 25. The compound of claim 24 wherein R_9 is fluoro or chloro.

26. The compound of claim 24 wherein R_{10} is methyl.

27. The compound of claim 25 wherein R_{10} is chloro.

25

28. The compound of claim 22 wherein R₉ is methyl.
29. The compound of claim 22 wherein R₉ and R₁₀ are not both H.
- 5 30. The compound of claim 22 wherein X is O.
31. The compound of claim 22 wherein X is N.
32. A composition comprising the compound of Claim 1 and a
10 pharmaceutically acceptable carrier.
33. The composition according to Claim 32 further comprising a compound having anti-HIV activity.
- 15 34. The composition according to Claim 33 wherein the compound having anti-HIV activity is effective to inhibit the function of a target selected from the group consisting of HIV protease, a nucleoside reverse transcriptase, a non-nucleoside reverse transcriptase or HIV entry.
- 20 35. The composition of claim 33 wherein the compound having anti-HIV activity is selected from:
- (a) an HIV protease inhibitor,
 - (b) a nucleoside reverse transcriptase inhibitor,
 - 25 (c) a non-nucleoside reverse transcriptase inhibitor,
 - (d) an HIV-entry inhibitor, and
 - (e) an immunomodulator,
- or combination thereof.
- 30 36. A method of inhibiting the function of the HIV integrase comprising contacting the HIV integrase with the compound of Claim 1.

37. A method of treating an HIV infection in a patient, comprising administering to the patient a therapeutically effective amount of the compound of Claim 1, or a pharmaceutically acceptable enantiomer, diastereomer, solvate, prodrug or salt thereof.

5

38. The method according to Claim 37 wherein the compound is effective to inhibit the function of the HIV integrase.

39. The method according to Claim 37 further comprising administering
10 another compound having anti-HIV activity prior to, after or simultaneously with the compound of Claim 1.

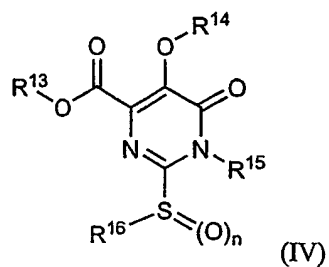
40. The method according to Claim 39 wherein the compound having anti-
HIV activity is effective to inhibit the function of a target selected from the group
15 consisting of HIV protease, a nucleoside reverse transcriptase, a non-nucleoside reverse transcriptase or HIV entry.

41. The method according to Claim 39 wherein the other compound having
anti-HIV activity is effective to inhibit the function of target in the HIV life cycle
20 other than the HIV integrase.

42. Use of the compound of Claim 1 for the manufacture of a medicament for treating HIV infection in a patient.

25 43. Use of the composition of Claim 32 for the manufacture of a medicament for treating HIV infection in a patient.

44. A compound of the formula:



wherein;

5

n is 0, 1 or 2;

R^{13} is -H, -C₁₋₆ alkyl, C₁₋₆ cycloalkyl, -aryl, C₁₋₆ alkyl-cycloalkyl, -C₁₋₆ alkyl-aryl, or -C₁₋₆ alkyl-diaryl,

10

R^{14} is -H, -C₁₋₆ alkyl, C₁₋₆ cycloalkyl, -aryl, C₁₋₆ alkyl-aryl, or -C₁₋₆ alkyl-diaryl,

R^{15} is -H, -C₁₋₆ alkyl, C₁₋₆ alkyl-cycloalkyl, -aryl, or -C₁₋₆ alkyl-aryl, and

15 R^{16} is -H, -C₁₋₆ alkyl, C₁₋₆ cycloalkyl, -aryl, C₁₋₆ alkyl-aryl, or -C₁₋₆ alkyl-diaryl.

(19) World Intellectual Property
Organization
International Bureau



(43) International Publication Date
29 July 2004 (29.07.2004)

PCT

(10) International Publication Number
WO 2004/062613 A3

(51) International Patent Classification⁷: **C07D 401/04**,
413/04, 417/04, A61K 31/506

(21) International Application Number:
PCT/US2004/000642

(22) International Filing Date: 12 January 2004 (12.01.2004)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
60/439,594 13 January 2003 (13.01.2003) US

(71) Applicant (for all designated States except US): **BRIS-
TOL-MYERS SQUIBB COMPANY** [US/US]; Route
206 And Provinceline Road, Princeton, NJ 08543-4000
(US).

(72) Inventors; and

(75) Inventors/Applicants (for US only): **WALKER, Michael**,
A. [US/US]; 25 Royal Oak Drive, Durham, CT 06422
(US). **GULGEZE, Hatice, Belgin** [US/US]; 623 Town
Colony Drive, Middletown, CT 06457 (US). **NAIDU,**
Narasimhulu, B. [IN/US]; 12 Skeetfield Point Road,
Durham, CT 06422 (US). **SORENSEN, Margaret, E.**
[US/US]; 27 Harrison Street, Apt. 13, Meriden, CT 06450
(US). **UEDA, Yasutsugu** [JP/US]; 46 Olde Orchard Road,
Clinton, CT 06413 (US). **MATISKELLA, John** [US/US];
130 High Hill Road, Wallingford, CT 06492 (US).

(74) Agents: **DUBOFF, Samuel et al.**; P.O. Box 4000, Prince-
ton, NJ 08543-4000 (US).

(81) Designated States (unless otherwise indicated, for every
kind of national protection available): AE, AG, AL, AM,
AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN,
CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI,
GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE,
KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD,
MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG,
PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM,
TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM,
ZW.

(84) Designated States (unless otherwise indicated, for every
kind of regional protection available): ARIPO (BW, GH,
GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW),
Eurasian (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), Euro-
pean (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR,
GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK,
TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW,
ML, MR, NE, SN, TD, TG).

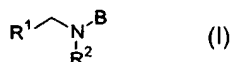
Published:

- with international search report
- before the expiration of the time limit for amending the
claims and to be republished in the event of receipt of
amendments

(88) Date of publication of the international search report:
4 November 2004

For two-letter codes and other abbreviations, refer to the "Guid-
ance Notes on Codes and Abbreviations" appearing at the begin-
ning of each regular issue of the PCT Gazette.

(54) Title: HIV INTEGRASE INHIBITORS



(57) Abstract: Compounds useful for treating HIV are disclosed having the general formula (I): wherein R¹, R², and B are as defined in the specification. Compositions containing the compounds and methods for using the compounds to inhibit HIV are also disclosed.

WO 2004/062613 A3

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US04/00642

A. CLASSIFICATION OF SUBJECT MATTER

IPC(7) : C07D 401/04, 413/04, 417/04; A61K 31/506

US CL : 544/3, 123, 319, 320, 321; 514/222.2, 235.8, 252.14, 272, 273

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 544/3, 123, 319, 320, 321; 514/222.2, 235.8, 252.14, 272, 273

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
CAS ONLINE**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 6,096,753 A (SPOHR et al) 01 August 2000 (01.08.2000), see entire document.	1-35

☐ Further documents are listed in the continuation of Box C.☐ See patent family annex.

* Special categories of cited documents:

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier application or patent published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

- "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
- "&" document member of the same patent family

Date of the actual completion of the international search

23 August 2004 (23.08.2004)

Date of mailing of the international search report

15 SEP 2004

Name and mailing address of the ISA/US

Mail Stop PCT, Attn: ISA/US
Commissioner for Patents
P.O. Box 1450
Alexandria, Virginia 22313-1450

Facsimile No. (703) 305-3230

Authorized officer

Deepak R. Rao

Telephone No. (571) 272-1600

**This Page is Inserted by IFW Indexing and Scanning
Operations and is not part of the Official Record**

BEST AVAILABLE IMAGES

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images include but are not limited to the items checked:

- ☒ **BLACK BORDERS**
- ☐ **IMAGE CUT OFF AT TOP, BOTTOM OR SIDES**
- ☐ **FADED TEXT OR DRAWING**
- ☐ **BLURRED OR ILLEGIBLE TEXT OR DRAWING**
- ☐ **SKEWED/SLANTED IMAGES**
- ☐ **COLOR OR BLACK AND WHITE PHOTOGRAPHS**
- ☐ **GRAY SCALE DOCUMENTS**
- ☐ **LINES OR MARKS ON ORIGINAL DOCUMENT**
- ☐ **REFERENCE(S) OR EXHIBIT(S) SUBMITTED ARE POOR QUALITY**
- ☐ **OTHER:** _____

IMAGES ARE BEST AVAILABLE COPY.

As rescanning these documents will not correct the image problems checked, please do not report these problems to the IFW Image Problem Mailbox.

END PAGE SLASH (USPTO)